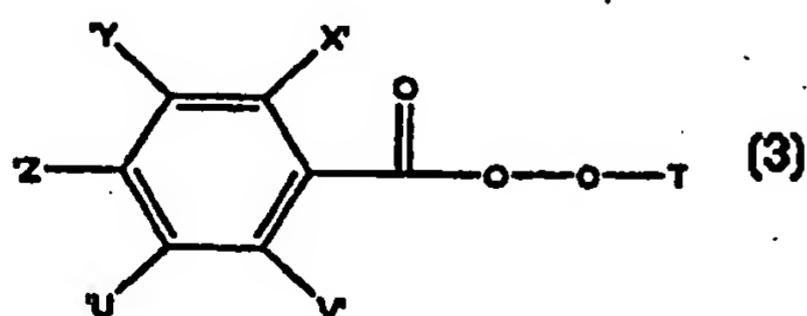
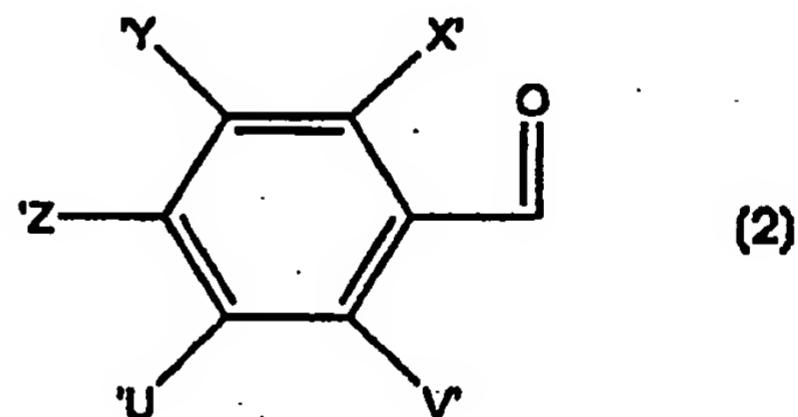
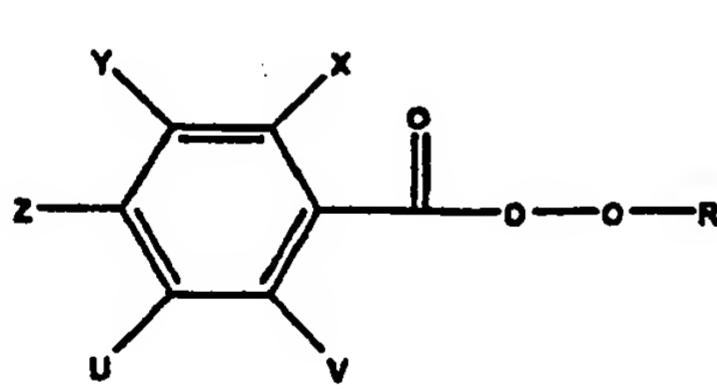




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(54) Title: PROCESS FOR INCREASING THE MELT STRENGTH OF POLYPROPYLENE



(57) Abstract

A process for modifying a polypropylene (co)polymer wherein said process comprises melt mixing the polypropylene (co)polymer in the presence of an initiator wherein said initiator is selected from the group defined by formula (1), wherein R is selected from the group consisting of optionally substituted C₁ to C₁₈ acyl, optionally substituted C₁ to C₁₈ alkyl, aroyl defined by formula (2), and compounds of formula (3), wherein U, V, X, Y, Z, U', V', X', Y' and Z' are independently selected from the group consisting hydrogen; halogen; C₁-C₁₈ alkyl; C₁-C₁₈ alkoxy, aryloxy, acyl, acyloxy, aryl, carboxy, alkoxycarbonyl, aryloxycarbonyl, trialkyl silyl, hydroxy, or a moiety of formula (4), and wherein T is alkylene.

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PROCESS FOR INCREASING THE MELT STRENGTH OF POLYPROPYLENE

- 5 The present invention relates to polypropylene homopolymers and copolymers. In particular, the present invention relates to a process for increasing the melt strength and/or the extensional melt viscosity of said polymers by melt phase processing.

The melt strength and extensional viscosity of linear or straight chain polymers, such as 10 polypropylene, decreases rapidly with temperature. By contrast, polymers such as low density polyethylene which are highly branched retain relatively high melt strengths and extensional viscosities. It is generally understood that the difference in melt strengths and extensional viscosities is attributable to the presence of long chain branching in polymers such as low density polyethylene. Long chain branching allows a greater degree of chain 15 entanglement.

A number of methods for increasing the melt strength/extensional viscosity of polypropylene and related polymers through the introduction of branching or a limited degree of crosslinking in a process involving reactive extrusion have been proposed and are summarised in a recent 20 paper by Wang et al. (Wang, X., Tzoganakis, C., and Rempel, G.L., *J. Appl. Polym. Sci.*, 1996, 61, 1395). One such process involves the reactive extrusion of polypropylene with a polyfunctional monomer/initiator combination. For example, the use of pentaerythritol triacrylate in combination with 2,5-dimethyl-2,5-di(t-butylperoxy)hexane (DHBP).

25 White (US 5578682) has disclosed the use of various polyunsaturated crosslinking agents (for example, bismaleimide derivatives) in combination with free radical initiators to achieve an increase in the melt strength various polymers.

It is well known that the melt phase processing of polypropylene leads to mechanochemical 30 degradation. The processing of polypropylene in the presence of free radical initiators

- 2 -

provides an increased rate of degradation. This controlled degradation of polypropylene is used commercially for the production of controlled rheology resins having reduced polydispersity and reduced die swell (Lambla, M. in *Comprehensive Polymer Science*, Pergamon, New York 1992, vol Suppl. 1, p 619; Hogt, A.H., Meijer, J., Jelinic, J. in 5 Reactive Modifiers for Polymers, Al-Malaika, S. Ed., Chapman & Hall, London, 1996, p 84.). The degradation of polypropylene as described therein results in a lowering of melt strength.

The batch modification of polypropylene to produce crosslinked (insoluble) polypropylene by 10 treatment with peroxides is described by Borsig et al. (Borsig, E., Fiedlerova, A., Lazar, M. J., *Macromol. Sci, Chem.*, 1981, A16, 513). Initiators which produce benzoyloxy radicals or phenyl radicals are described as being more efficient in inducing crosslinking or grafting than those which produce t-butoxy or alkyl radicals. The process requires the use of high levels of peroxide. The use of polyfunctional monomers as coagents to retard degradation and 15 enhance crosslinking is described by Chodak, I.; Fabianova, K.; Borsig, E.; Lazar, M. Agnew. *Makromol. Chem.*, 1978, 69, 107.

DeNicola (EP 384331A2) has disclosed a means to produce a branched propylene polymer material showing a nett increase in the weight average molecular weight by solid state 20 modification of predominantly isotactic semi-crystalline linear polypropylene. The process described in EP384331A2 involves blending peroxides with short half lives (eg peroxy dicarbonates) with linear propylene polymer in a mixing vessel at temperatures from 23°C to 120°C in an inert atmosphere and continuing to mix for a period of time until the peroxide decomposes and polymer fragmentation and branching occurs without significant gelation of 25 the polymer. DeNicola states that at temperatures greater than 120°C no branching or melt strength enhancement is achieved.

U.S. 5,464,907 teaches that certain unsaturated maleate or itaconate derived peroxides may be used to induce grafting in polypropylene and α -olefin copolymers. They report that use

- 3 -

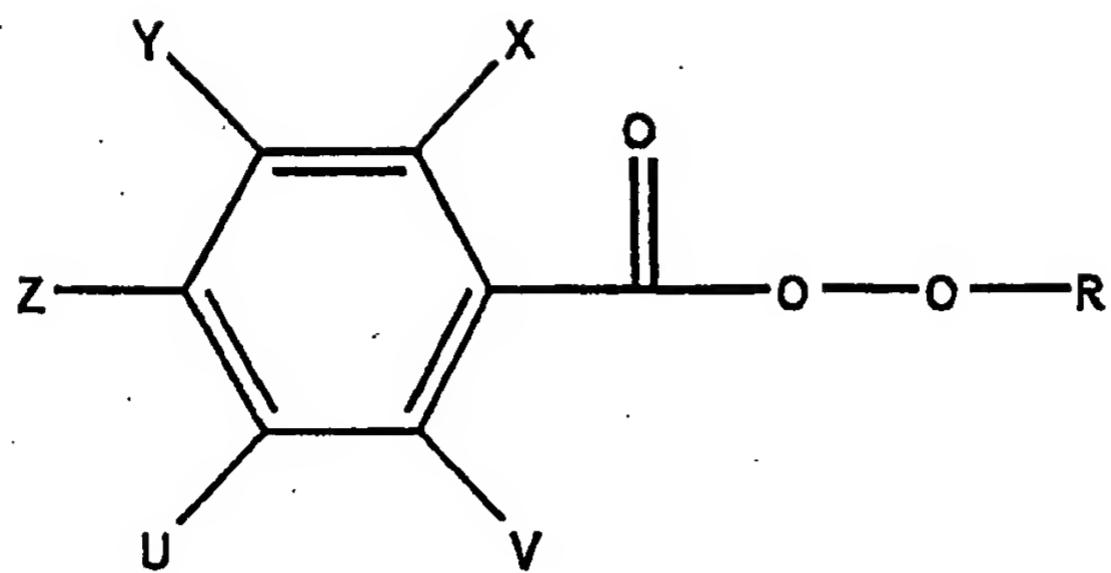
of other peroxides generally results in chain degradation.

- Polypropylene is also known to undergo substantial degradation during melt phase grafting of monofunctional monomers, for example maleic anhydride and glycidyl methacrylate. It
5 has also been reported that the degradation that accompanies grafting of these monomers to polypropylene may be reduced by the addition of relatively high concentrations of certain comonomers including styrene (see, for example Sun, Y.-J., Hu, G.-H., and Lambla, M., *Angew. Makromol. Chem.*, 1995, 229, 1; Chen, L-F., Wong, B. and Baker, W.E. *Polym. Eng. Sci.* 1996, 36, 1594.) Sun et al. report that there is degradation (as indicated by an
10 overall decrease in molecular weight) when styrene alone is grafted onto polypropylene even when a relatively high concentration is used (4 moles/100g PP). Either 2,5-dimethyl-2,5-(t-butylperoxy)hex-3-yne or 2,5-dimethyl-2,5(t-butylperoxy)-hexane(DHBP) was used as the initiator in these experiments.

15 We have found that melt mixing polypropylene homopolymer or ethylene-polypropylene copolymer in the presence of a suitable initiator provides one or more of the following: increased melt strength; increased extensional viscosity; increased molecular weight; and broadened molecular weight distribution.

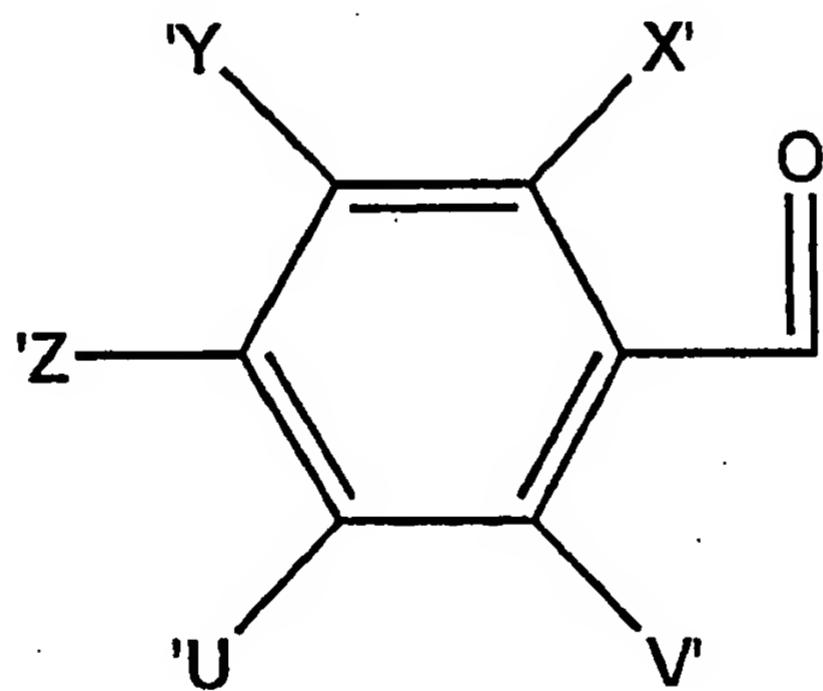
20 According to the present invention there is provided a process for modifying a polypropylene (co)polymer wherein said process comprises melt mixing the polypropylene (co)polymer in the presence of an initiator wherein said initiator is selected from the group defined by formula 1.

- 4 -



Formula 1

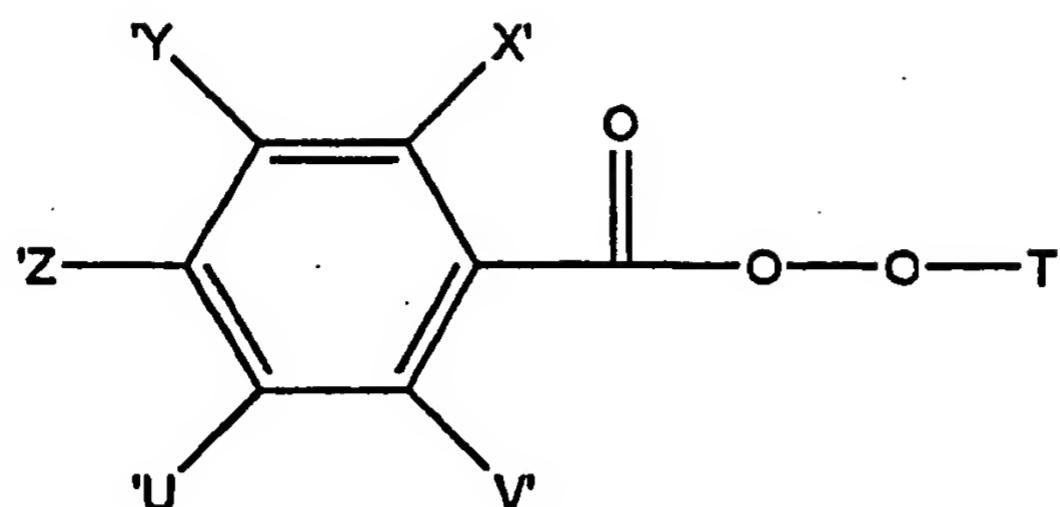
wherein R is selected from the group consisting of optionally substituted C₁ to C₁₈ acyl,
5 optionally substituted C₁ to C₁₈ alkyl, aroyl defined by formula 2,



Formula 2

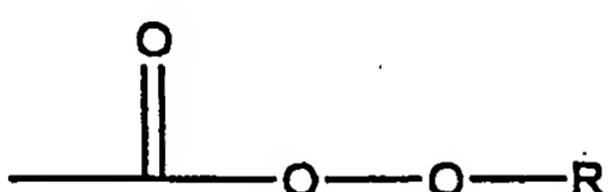
10 and compounds of formula 3,

- 5 -



Formula 3

wherein U, V, X, Y, Z, U', V', X', Y' and Z' are independently selected from the group
 5 consisting hydrogen; halogen; C₁-C₁₈ alkyl; C₁-C₁₈ alkoxy, aryloxy, acyl, acyloxy, aryl,
 carboxy, alkoxycarbonyl, aryloxycarbonyl, trialkyl silyl, hydroxy,
 or a moiety of formula 4,



10

Formula 4

and wherein T is alkylene.

Advantageously the thus formed modified polypropylene may be obtained without the associated production of significant and detrimental amounts of gels.

15

Polymers suitable for use in the present invention include a wide variety of polypropylene homopolymers, copolymers and blends containing one or more polypropylene homopolymers and/or copolymers.

20 Suitable polypropylene homopolymers include isotactic polypropylene, atactic polypropylene and syndiotactic polypropylene. Commercial isotactic polypropylene having a proportion of

- 6 -

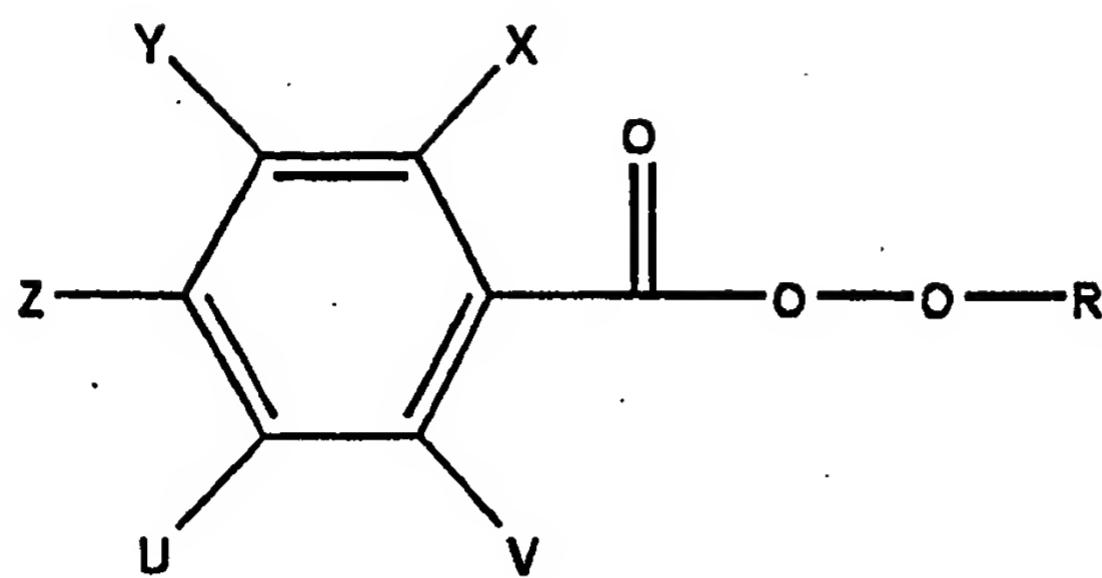
meso/dyads of greater than 90% is preferably used in the process of the present invention. Isotactic polypropylene is a semi-crystalline polymer having a number of properties which have made it one of the most widely used commercial polymers. These properties include heat resistance, stress cracking resistance, chemical resistance, toughness, and low 5 manufacturing costs. However, the melt strength of isotactic polypropylene as measured directly by extensional viscosity or use of a commercial melt strength tester or indirectly by more qualitative measures such as drop time or die swell ratio is relatively low. This relatively low melt strength limits the use of polypropylene in applications such as foam extrusion, thermoforming and film blowing. In order to use polypropylene in such 10 applications it is necessary to employ sophisticated processing equipment. The present invention now permits this already widely used commercial polymer to be used in an even wider range of applications.

Polypropylene copolymers include copolymers of propylene and other monomers with such 15 other monomers being present preferably in amounts of up to 10%wt/wt. A preferred comonomer is ethylene.

The present invention is also applicable to other polymers comprising α -olefin monomers. It is preferable that any such α -olefins are present in the polymer to be modified in amounts 20 in excess of 90%wt/wt. α -olefins include propene, 1-butene, 1-pentene and 1-hexene.

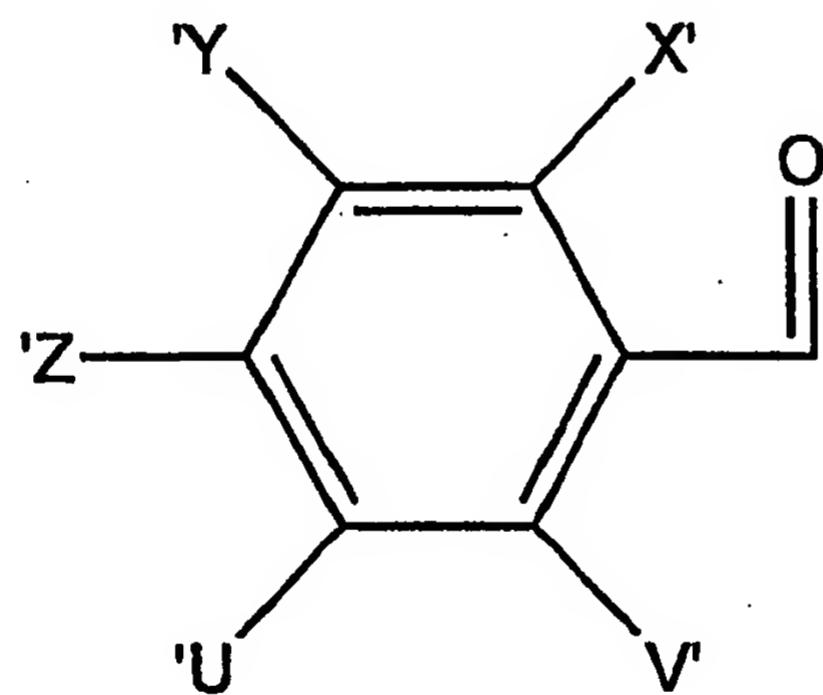
The initiators for use in the present invention may be selected from the group defined by formula 1.

- 7 -



Formula 1

5 wherein R is selected from the group consisting of optionally substituted C₁ to C₁₈ acyl, optionally substituted C₁ to C₁₈ alkyl, aroyl defined by formula 2,

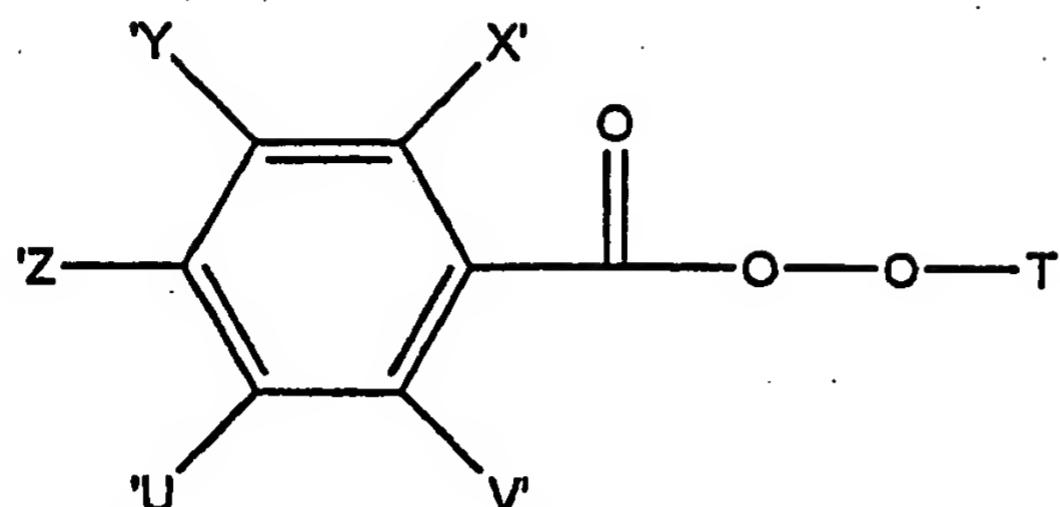


Formula 2

10

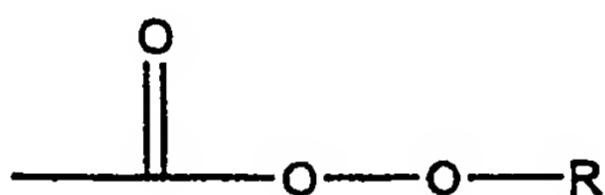
and compounds of formula 3,

- 8 -



Formula 3

wherein U, V, X, Y, Z, U', V', X', Y' and Z' are independently selected from the group 5 consisting hydrogen; halogen; C1-C18 alkyl; C1-C18 alkoxy, aryloxy, acyl, acyloxy, aryl, carboxy, alkoxycarbonyl, aryloxycarbonyl, trialkyl silyl, hydroxy, or a moiety of formula 4,



10

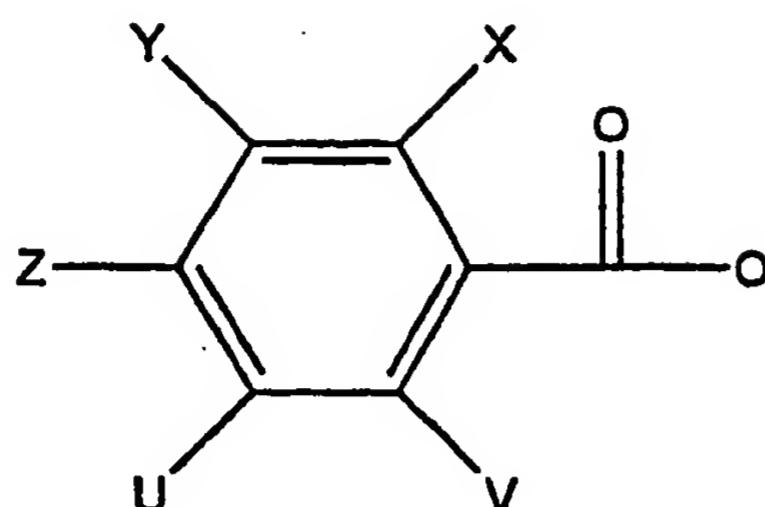
Formula 4

and wherein T is alkylene.

The alkyl, including acyl and alkoxy, groups included in the initiators of formula 1 may 15 include hetero atoms within the carbon chain (eg polyalkylene oxide) and may be branched or unbranched and may be substituted with one or more groups such as with alkyl, aryl, alkoxy or halogen substituents.

Without wishing to be bound by theory, it is believed that the aroyloxy radical of formula 5
20

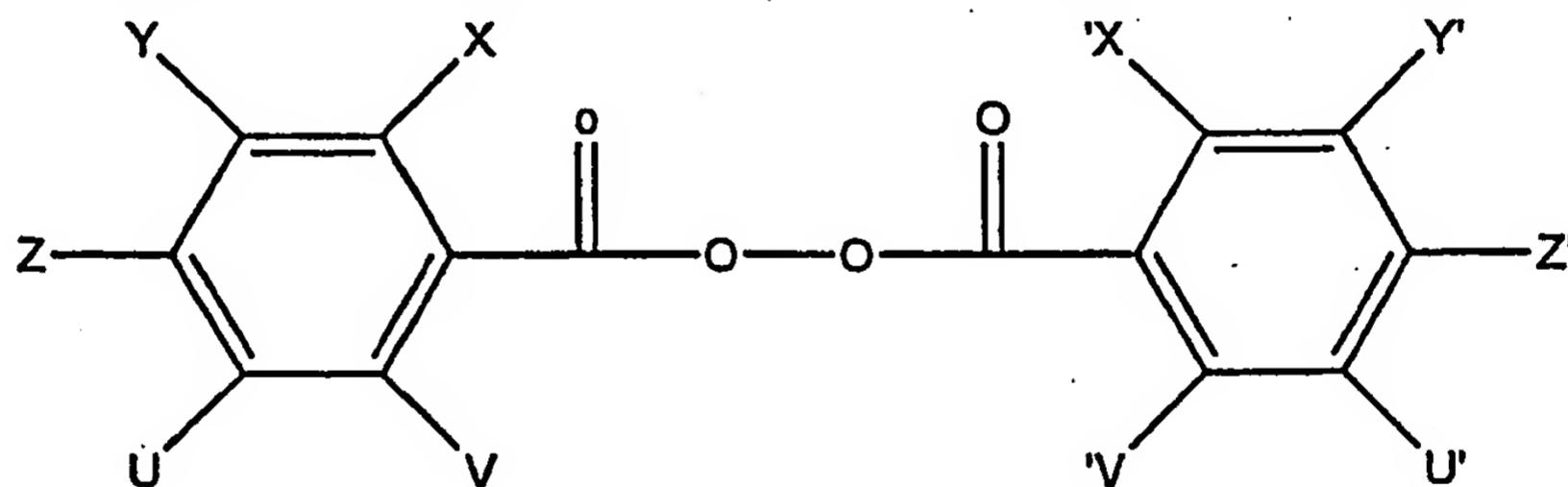
- 9 -



Formula 5

, where U, V, X, Y and Z are as hereinabove defined, provide the surprising increase in melt strength. Other compounds which generate these aroyloxy radicals may also be used in the present invention.

A preferred class of initiators of formula 1 are diaryl peroxides of formula 6.



10

Formula 6

where X, Y, Z, U, V, X', Y', Z', U', V' are independently selected from the group consisting of hydrogen and C₁ - C₁₈ alkyl where at least one of X, Y, Z, U, V and X', Y', Z', U', V' are not hydrogen.

15

Diaryl peroxides of formula 6 include Dibenzoyl peroxide, o,o'-Bis(methylbenzoyl) peroxide, p,p'-Bis(methylbenzoyl) peroxide, M,M'-Bis(methylbenzoyl) peroxide, o,m'-Bis(methylbenzoyl)

- 10 -

- peroxide, o,p'-Bis(methylbenzoyl) peroxide, m,p'-Bis(methylbenzoyl) peroxide, Bis(ethylbenzoyl) peroxide (all isomers), Bis(propylbenzoyl) peroxide (all isomers), Bis(butylbenzoyl) peroxide (all isomers), Bis(pentylbenzoyl) peroxide (all isomers), Bis(hexylbenzoyl) peroxide (all isomers), Bis(heptylbenzoyl) peroxide (all isomers), Bis(octylbenzoyl) peroxide (all isomers),
5 Bis(nonylbenzoyl) peroxide (all isomers), Bis(methoxybenzoyl) peroxide (all isomers), Bis(ethoxybenzoyl) peroxide (all isomers), Bis(propoxybenzoyl) peroxide (all isomers), Bis(butoxybenzoyl) peroxide (all isomers), Bis(pentoxybenzoyl) peroxide (all isomers), Bis(hexyloxybenzoyl) peroxide (all isomers), Bis(heptyloxybenzoyl) peroxide (all isomers), Bis(octyloxybenzoyl) peroxide (all isomers), Bis(nonyloxybenzoyl) peroxide (all isomers),
10 Bis(chlorobenzoyl) peroxide (all isomers), Bis(fluorobenzoyl) peroxide (all isomers), Bis(bromobenzoyl) peroxide (all isomers), Bis(dimethylbenzoyl) peroxide (all isomers), Bis(trimethylbenzoyl) peroxide (all isomers), Bis(tert-butylbenzoyl)peroxide (all isomers), Bis(di-tert-butylbenzoyl)peroxide (all isomers), Bis(tertbutoxybenzoyl)peroxide (all isomers), Bis(ditrimethylsilylbenzoyl) peroxide (all isomers), Bis(heptafluoropropylbenzoyl) peroxide (all
15 isomers), Bis(2,6-dimethyl-4- trimethylsilyl benzoyl) peroxide and isomers, 2,2'(dioxydicarbonyl) bis - Benzoic acid dibutyl ester where the term "all isomers" refers to any variation in the position of the ring substituent as well as the structure of the substituent itself i.e. for propyl; n-propyl and isopropyl.

20 Examples of aromatic peresters of formula 1 include the following: tert-butyl perbenzoate, tert-butyl (methyl)perbenzoate (all isomers), tert-butyl (ethyl)perbenzoate (all isomers), tert-butyl (octyl)perbenzoate (all isomers), tert-butyl (nonyl)perbenzoate (all isomers), tert-amyl perbenzoate, tert-amyl (methyl)perbenzoate (all isomers), tert-amyl (ethyl)perbenzoate (all isomers), tert-amyl (octyl)perbenzoate (all isomers), tert-amyl (nonyl)perbenzoate (all isomers),
25 tert-amyl (methoxy)perbenzoate (all isomers), tert-amyl (octyloxy)perbenzoate (all isomers), tert-amyl (nonyloxy)perbenzoate (all isomers), 2-ethylhexyl perbenzoate, 2-ethylhexyl (methyl)perbenzoate (all isomers), , 2-ethylhexyl (ethyl)perbenzoate (all isomers), 2-ethylhexyl (octyl)perbenzoate (all isomers), , 2-ethylhexyl (nonyl)perbenzoate (all isomers), 2-ethylhexyl (methoxy)perbenzoate (all isomers), 2-ethylhexyl (ethoxy)perbenzoate (all isomers), 2-
30 ethylhexyl (octyloxy)perbenzoate (all isomers), 2-ethylhexyl (nonyloxy)perbenzoate (all isomers)

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The initiators for use in the present invention also include compounds of formula 1 where at least one of U, V, X, Y, Z, U', V', X' Y' and Z' is a moiety of formula 4 where R is as defined above. Preferably there is no more than one moiety of formula 4 per aromatic ring. Such initiators are di or higher functional peroxides and may include polymeric peroxides 5 such as Bis (tertbutylmonoperoxy phthaloyl) diperoxy terephthalate, Bis (tertamylmonoperoxy phthaloyl) diperoxy terephthalate diacetyl phthaloyl diperoxide, dibenzoyl phthaloyl diperoxide, bis(4 methylbenzoyl) phthaloyl diperoxide, diacetyl terephthaloyl di peroxide, dibenzoyl terephthaloyl diperoxide, Poly[dioxycarbonyldioxy(1,1,4,4-tetramethyl-1,4-butanediyl)] peroxide.

10

It is described that the initiators are selected such that it has an appropriate decomposition temperature (half life), solubility, and reactivity and such that the groups R, T, X, Y, Z, U, V, X', Y', Z', U', V' give no adverse reaction under the conditions of the process. Preferred 15 peroxides will have a 0.1 hour half life in the range 100 - 170°C.

The amount of initiator used in the process of the present invention should be an effective amount to achieve the desired increase in melt strength. Melt strength is considered in the art to be an indication of long-chain branching in polyolefins. It is preferable in the process 20 of the present invention that long-chain branching predominates over crosslinking in the reaction between the initiator and the polypropylene (co)polymer. Crosslinking of the polypropylene (co)polymer may result in the formation of gels which disrupt the appearance of the polypropylene (co)polymer. In the process of the present invention it is desirable to control the degree and distribution of crosslinking and keep the level of crosslinking as 25 uniform and as low as necessary to produce the desired effects. The amount of crosslinking which occurs in the polypropylene (co)polymer is dependant upon the amount of initiator melt mixed with the polypropylene (co)polymer. The amount of crosslinking is also dependent upon the degree of mixing as any regions high in initiator concentration will result in excessive localised crosslinking and the formation of gels. It is desirable that good 30 distributive and dispersive mixing be employed to promote even distribution of the initiator

- 12 -

in the polypropylene (co)polymer so as to minimise the variation in initiator concentration throughout the polypropylene (co)polymer and reduce the likelihood of the formation of gels.

Preferably the initiator will be present in the range of from 0.004 to 0.25 moles of initiator per kg of the polypropylene homopolymer or copolymer (polypropylene (co)polymer). The more preferred range being from 0.006 to 0.10 moles of initiator per kg. of the polypropylene (co)polymer and even more preferred range being from 0.01 to 0.05 moles of initiator per kg of the polypropylene (co)polymer.

10 The initiator is preferably introduced into the polymer melt directly, either neat (as a powder or a liquid), dispersed or dissolved in a suitable medium (for example, dissolved in 2-butanone) or adsorbed on polymer pellets or powder which are added as a masterbatch. It is desirable that the initiator is rapidly mixed with the polymer melt at a rate in keeping with the half life of the initiator at the processing temperature of the polypropylene (co)polymer.

15

The initiator may be added either alone, or along with the polypropylene (co)polymer, or with any other polymer, additive or filler, so that the polymer melts and mixes with the initiator as it is decomposing. When the initiator is fed to the main feed throat of the extruder it is preferred to have a barrel temperature which is relatively low in the region adjacent to the 20 main feed throat and increases towards the die to prevent premature decomposition of the peroxide.

Preferably the initiators for use in the present invention are selected from the group consisting of Dibenzoyl peroxide, o,o'-Bis(methylbenzoyl) peroxide, p,p'-Bis(methylbenzoyl) peroxide, o,o'-
25 Bis(methylbenzoyl) peroxide, o,m'-Bis(methylbenzoyl) peroxide, o,p'-Bis(methylbenzoyl) peroxide, m,p'-Bis(methylbenzoyl) peroxide, Bis(ethylbenzoyl) peroxide (all isomers), Bis(propylbenzoyl) peroxide (all isomers), Bis(butylbenzoyl) peroxide (all isomers), Bis(pentylbenzoyl) peroxide (all isomers), Bis(hexylbenzoyl) peroxide (all isomers), Bis(heptylbenzoyl) peroxide (all isomers), Bis(octylbenzoyl) peroxide (all isomers),
30 Bis(nonylbenzoyl) peroxide (all isomers), Bis(methoxybenzoyl) peroxide (all isomers),

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Bis(ethoxybenzoyl) peroxide (all isomers), Bis(propoxybenzoyl) peroxide (all isomers), Bis(butoxybenzoyl) peroxide (all isomers), Bis(pentoxybenzoyl) peroxide (all isomers), Bis(hexyloxybenzoyl) peroxide (all isomers), Bis(heptyloxybenzoyl) peroxide (all isomers), Bis(octyloxybenzoyl) peroxide (all isomers), Bis(nonyloxybenzoyl) peroxide (all isomers),
5 Bis(chlorobenzoyl) peroxide (all isomers), Bis(fluorobenzoyl) peroxide (all isomers), Bis(bromobenzoyl) peroxide (all isomers), Bis(dimethylbenzoyl) peroxide (all isomers), Bis(trimethylbenzoyl) peroxide (all isomers), Bis(tert-butylbenzoyl)peroxide (all isomers), Bis(di-tert-butylbenzoyl)peroxide (all isomers), Bis(tertbutoxybenzoyl)peroxide (all isomers), Bis(dtrimethylsilylbenzoyl) peroxide (all isomers), Bis(heptafluoropropylbenzoyl) peroxide (all
10 isomers), Bis(2,4-dimethyl-6- trimethylsilyl benzoyl) peroxide and isomers tert-amyl perbenzoate, tert-amyl (methyl)perbenzoate (all isomers), tert-amyl (ethyl)perbenzoate (all isomers), tert-amyl (octyl)perbenzoate (all isomers), tert-amyl (nonyl)perbenzoate (all isomers), tert-amyl (methoxy)perbenzoate (all isomers), tert-amyl (octyloxy)perbenzoate (all isomers), tert-amyl (nonyloxy)perbenzoate (all isomers), Bis (tertamylmonoperoxy phthaloyl) diperoxy terephthalate,
15 diacetyl phthaloyl diperoxide, dibenzoyl phthaloyl diperoxide, bis(4-methylbenzoyl) phthaloyl diperoxide, diacetyl terephthaloyl di peroxide and dibenzoyl terephthaloyl diperoxide.

More preferably the initiators are selected from the group consisting of dibenzoyl peroxide, o,o'-Bis(methylbenzoyl) peroxide, p,p'-Bis(methylbenzoyl) peroxide, M,M'-Bis(methylbenzoyl)
20 peroxide, o,m'-Bis(methylbenzoyl) peroxide, o,p'-Bis(methylbenzoyl) peroxide, m,p'-Bis(methylbenzoyl) peroxide.

The initiators may optionally be used in combination with one or more monomers.

25 Preferably the one or more monomers are selected from the group consisting of monoene monomer. It will be understood by those skilled in the art that by the term "monoene monomer" it is meant a monomer having a single reactive double bond.

The preferred monoene monomer(s) or mixtures thereof include vinyl monomers of structure
30 $\text{CH}_2 = \text{CHX}$ where X is chosen so as to confer the desired reactivity and solubility.

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More preferred monomers include styrene. The amount of monomer will preferably be up to 5 times the total moles of initiator added to the polypropylene (co)polymer. The most preferred range being 1 to 4 times the total moles of initiator added to the polypropylene (co)polymer.

5

The monomer may be added with the polypropylene (co)polymer or it can be added prior to the initiator, with the initiator or subsequent to the initiator. However it is preferred to have the monomer mixed and dispersed into the polymer melt before the initiator has substantially decomposed. The monomer is preferably introduced into the polymer melt directly, either 10 neat (as a powder or a liquid), dispersed or dissolved in a suitable medium (for example, dissolved in 2-butanone) or adsorbed on polymer pellets or powder which are added as a masterbatch.

Preferred initiators for use in combination with monomers include Dibenzoyl peroxide, o,o'-
15 Bis(methylbenzoyl) peroxide, p,p'-Bis(methylbenzoyl) peroxide, M,M'-Bis(methylbenzoyl)
peroxide, o,m'-Bis(methylbenzoyl) peroxide, o,p'-Bis(methylbenzoyl) peroxide, m,p'-
Bis(methylbenzoyl) peroxide, Bis(ethylbenzoyl) peroxide (all isomers), Bis(propylbenzoyl)
peroxide (all isomers), Bis(butylbenzoyl) peroxide (all isomers), Bis(pentylbenzoyl) peroxide (all
isomers), Bis(hexylbenzoyl) peroxide (all isomers), Bis(heptylbenzoyl) peroxide (all isomers),
20 Bis(octylbenzoyl) peroxide (all isomers), Bis(nonylbenzoyl) peroxide (all isomers),
Bis(methoxybenzoyl) peroxide (all isomers), Bis(ethoxybenzoyl) peroxide (all isomers),
Bis(propoxybenzoyl) peroxide (all isomers), Bis(butoxybenzoyl) peroxide (all isomers),
Bis(pentoxybenzoyl) peroxide (all isomers), Bis(hexyloxybenzoyl) peroxide (all isomers),
Bis(heptyloxybenzoyl) peroxide (all isomers), Bis(octyloxybenzoyl) peroxide (all isomers),
25 Bis(nonyloxybenzoyl) peroxide (all isomers), Bis(chlorobenzoyl) peroxide (all isomers),
Bis(fluorobenzoyl) peroxide (all isomers), Bis(bromobenzoyl) peroxide (all isomers),
Bis(dimethylbenzoyl) peroxide (all isomers), Bis(trimethylbenzoyl) peroxide (all isomers),
Bis(tert-butylbenzoyl) peroxide (all isomers), Bis(di-tert-butylbenzoyl) peroxide (all isomers),
Bis(tert-butoxybenzoyl) peroxide (all isomers), Bis(ditrimethylsilylbenzoyl) peroxide (all isomers),
30 Bis(heptafluoropropylbenzoyl) peroxide (all isomers), Bis(2,4-dimethyl-6-trimethylsilyl benzoyl)

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peroxide and isomers, 2,2'(dioxydicarbonyl) bis - Benzoic acid dibutyl ester, tert-butyl perbenzoate, tert-butyl (methyl)perbenzoate (all isomers), tert-butyl (ethyl)perbenzoate (all isomers), tert-butyl (octyl)perbenzoate (all isomers), tert-butyl (nonyl)perbenzoate (all isomers), tert-amyl perbenzoate, tert-amyl (methyl)perbenzoate (all isomers), tert-amyl (ethyl)perbenzoate
5 (all isomers), tert-amyl (octyl)perbenzoate (all isomers), tert-amyl (nonyl)perbenzoate (all isomers), tert-amyl (methoxy)perbenzoate (all isomers), tert-amyl (octyloxy)perbenzoate (all isomers), tert-amyl (nonyloxy)perbenzoate (all isomers), 2-ethylhexyl perbenzoate, 2-ethylhexyl (methyl)perbenzoate (all isomers), , 2-ethylhexyl (ethyl)perbenzoate (all isomers), 2-ethylhexyl (octyl)perbenzoate (all isomers), , 2-ethylhexyl (nonyl)perbenzoate (all isomers), 2-ethylhexyl
10 (methoxy)perbenzoate (all isomers), 2-ethylhexyl (ethoxy)perbenzoate (all isomers), 2-ethylhexyl (octyloxy)perbenzoate (all isomers), 2-ethylhexyl (nonyloxy)perbenzoate (all isomers), Bis (tertbutylmonoperoxy phthaloyl) diperoxy terephthalate, Bis (tertamylmonoperoxy phthaloyl)
diperoxy terephthalate diacetyl phthaloyl diperoxide, dibenzoyl phthaloyl diperoxide, bis(4
methylbenzoyl) phthaloyl diperoxide, diacetyl terephthaloyl di peroxide, dibenzoyl terephthaloyl
15 diperoxide and Poly[dioxycarbonyldioxy(1,1,4,4-tetramethyl-1,4-butanediyl)] peroxide.

Advantageously initiators may be selected to avoid undesirable by-products. In certain applications, it may be desirable to avoid the use of initiators which generate benzene. For example di toluoyl peroxides (bis methyl benzoyl peroxides) may be used in preference to
20 dibenzoyl peroxide.

The processability and other properties of the product may be improved by a chain scission step following the initial polymer modification step. This may be carried out by:
a) adding one or more additional initiators with or subsequent to the first initiator addition;
25 b) the use of high shear mixing;
c) the use of high temperatures;
d) the use combination is of one or more of (a) - (c) above.

This additional step in the production of a polymer enables tailoring the properties of the
30 product to meet the requirements of the desired application. For example, by this two stage

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process it is possible to produce materials with similar melt viscosity to the base polymer but a substantially increased melt strength. Use of the single stage process generally provides both an increase in melt strength and an increase in melt viscosity (see examples)

- 5 One or more additional initiators may be added to the polypropylene (co)polymer during the modification process either with or subsequent to the initiator and monomer addition. The additional initiator is typically added to give chain scission of the polypropylene (co)polymer so as to decrease the melt viscosity and improve the processability of the modified polypropylene (co)polymer. The additional initiator should be introduced to the polymer melt
- 10 after the first initiator or have a sufficiently long half-life relative to the first initiator such that its decomposition can be staged to occur after the initial polymer modification process. In some instances a polypropylene (co)polymer modified in accordance with the present invention may have a MFI < 1 g/10 min. With use of the additional initiator an MFI > 1 g/10 min may be achieved. The additional initiator may be selected from the group consisting of
- 15 2,5-dimethyl-2,5-di(t-butylperoxy)hexane (DHBP), dicumyl peroxide (DCP), t-butyl peroxy-2-ethylhexonate(TBEH), and dilauryl peroxide (DLP) or any other peroxide which may result in the overall chain scission of the polypropylene (co)polymer during melt processing. For example in the absence of the monoene monomers, t-butyl peroxybenzoate or other non-preferred initiators for use in the presence of the monomer may be preferably added as the
- 20 additional initiator. While the improvement in processability through chain scission normally results in some decrease in the melt strength/extensional viscosity of the modified polypropylene (co)polymer, the melt strength/extensional viscosity may still be acceptable, and improved over the unmodified polypropylene (co)polymer.
- 25 It is possible to combine the process of the present invention with other processes of polymer modification or with, for example, the addition of fillers, additives or stabilisers, or blending with other polymers. [which do not substantially interfere with the improved properties afforded by the process of the present invention].

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In the process of the present invention the polypropylene (co)polymer is melt mixed in the presence of initiator and monomer. Melt mixing may be carried out by any convenient means capable of mixing the polypropylene (co)polymer at temperatures above the melting point of the polypropylene (co)polymer.

5

Suitable apparatus for melt mixing the polypropylene (co)polymer include continuous and batch mixers. Suitable mixing equipment includes extruders such as single screw and twin screw extruders, static mixers, cavity transfer mixers and combinations of two or more thereof. It is preferred that the melt mixing is conducted in either a co- or counter- rotating 10 twin screw extruder.

The barrel set temperatures are preferably in the range 80-280 °C. Typical melt temperatures are in the range 170-290 °C.

15 In order to optimise the melt strength/extensional viscosity, the preferred melt temperatures are in the range 160 °C to 220 °C. This range provides optimal properties whilst minimising the amount of chain scission which occurs during processing. However, in some cases it may be desirable to use higher temperatures such as in the venting/discharge sections of single screw or twin screw extruders or to induce some chain scission in order to decrease the 20 molecular weight of the modified polypropylene (co)polymer and improve the processability of the modified polypropylene (co)polymer.

Typically, the die temperatures are in the range 180-290 °C.

25 Preferably the extrusion conditions are adjusted so that the polypropylene (co)polymer, initiator/monomer mixture are conveyed as quickly as possible into the melting/mixing zone to maximise the melt phase reaction (eg for twin screw extruders - high throughput rates, higher screw speeds under starve fed conditions). It is more preferred that the additives are added to and mixed with molten polypropylene (co)polymer to further enhance the melt phase

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reaction. Preferably residence times in the range of from 10 seconds to 5 minutes are selected depending upon the temperature profile, throughput rate and initiator levels. More preferred residence times are in the range of from 15 seconds to 120 seconds.

5 Vacuum venting can be applied to remove volatile by-products, solvents and/or excess monomer.

While not wishing to be limited by theory, it is believed that the effectiveness of the present invention is determined by three factors:

10

(a) The rate and specificity of the reaction of the aroyloxy or the derived phenyl radicals or substituted phenyl radicals with polypropylene, and the monomer if present. It is believed that the aroyloxy, phenyl or substituted aroyloxy or phenyl radicals show less specificity for abstraction of tertiary vs. secondary or primary hydrogens than do, for example, alkoxy or 15 alkyl radicals.

(b) The initiator half-life. Use of an initiator with a short initiator half-life will generate a locally high concentration of radicals thus increasing the likelihood of radical combination events.

20

(c) The solubility characteristics of the initiator in the polymer melt.

Without wishing to be bound by theory, peroxides that generate aroyloxy or aryl radicals (for example benzoyloxy, p-toluouloxy) are preferred over those that generate alkoxy radicals (for 25 example, t-butoxy radical, cumyloxy radical). It is believed and supported in the literature that the latter class of peroxides promote chain scission under the melt mixing conditions. While not wishing to be bound by the mechanism, it is believed that this effect is due to the specificity shown by the alkoxy radicals as opposed to the aroyloxy or aryl radicals generated by the peroxides of structure 1. Furthermore we believe that peroxides which generate both

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alkoxy and aroyloxy or aryl radicals (for example, t-butyl perbenzoate) show intermediate behaviour. It is believed that they promote less chain scission than peroxides which generate only alkoxy radicals (for example, dialkyl peroxides) when used alone and can be used to advantage in systems where a monomer coagent is employed. Preferred peresters are thus 5 those which generate alkoxy radicals which are not active in hydrogen abstraction (for example t-amyl perbenzoate).

Similarly, it is believed, without wishing to be bound by theory, that the effectiveness of the monomer is determined by:

10

(a) The solubility of the monomer in the polymer melt. For example, styrene is known to be soluble in molten polypropylene.

(b) The reactivity of the monomer towards polypropylene derived radicals.

15

(c) The propensity for the radical formed by addition of monomer to give combination or addition (which leads to branch or crosslink formation) vs. disproportionation or hydrogen abstraction. It is known that the benzylic radicals give predominantly combination and have low (with relation to other radicals) tendency to abstract hydrogen.

20

Other initiators and monomers that meet the above criteria may also be used to advantage in the present invention.

Surprisingly, the process of the present invention results in a polypropylene (co)polymer with 25 substantially increased melt strength. We have found that it is possible with the present invention to obtain a polypropylene (co)polymer which has a melt strength at least 25% greater than the melt strength of the base polymer. We have also found that it is possible to obtain an increase in melt strength of greater than 100% for a number of the polypropylene (co)polymers produced in accordance with the process of the present invention. Increases in

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melt strength were assessed using a Gottfert-Rheotens melt strength tester operated with a roller acceleration of 1.2 cm/sec² measuring the melt strength of a 2 mm strand of molten polypropylene (co)polymer (melt temperature of 210°C) which is fed to the Gottfert tester at ~ 4 g/min.

5

In a further aspect of the present invention there is provided a modified polypropylene (co)polymer produced according to the process described herein, wherein said modified polypropylene (co)polymer preferably has a melt strength at least 25 %, and more preferably at least 100 %, greater than the unmodified polypropylene (co)polymer.

10

The polypropylene (co)polymers produced according to the process of the present invention also may provide a significant increase in long-chain branching. Long-chain branching may be assessed by the Dow Rheology Index. Advantageously, the modified polypropylene (co)polymers may demonstrate a Dow Rheology Index (DRI) of greater than 1, preferably 15 at least 2 and most preferably greater than 50.

The process of the present invention may also be used to increase the melt elasticity of a polypropylene (co)polymer.

20 Advantageously, the process of the present invention also provides a means to alter the molecular weight, molecular weight distribution and/or degree and length of branching of polypropylene, ethylene-propylene copolymers, and analogous α -olefin copolymers with or without altering the melt strength of said polymers by melt processing.

25 The process of the present invention may provide a means to generally increase the molecular weight and broaden the molecular weight distribution and/or introduce branching of the polypropylene (co)polymer. This will not always equate to significant increases in the melt strength or extensional viscosity of the polymer that is being modified eg modification of a lower molecular weight polymer to broaden the molecular weight and/or induce shorter

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branches. Such a product may not necessarily demonstrate a high melt strength, but may demonstrate other desirable properties, for example improved filler uptake, mechanical properties, surface properties, thermal and morphological properties.

- 5 The modified polypropylene (co)polymer produced by the process of the present invention may be used either neat or blended with another polymer or other additives to provide the desired balance of properties in the polymer blend.

The modified polypropylene (co)polymers and blends may be used in a wide variety of 10 applications including thermoforming, blow moulding, tube or pipe extrusion, blown films, foams and extrusion coating.

The present invention may also be used in the recycling of waste polypropylene or materials containing waste polypropylene.

15

The increased melt strength of the modified polypropylene (co)polymers renders these (co)polymers more suitable for use in thermoforming applications. The modified polypropylene (co)polymers may be used to thermoform containers such as margarine tubs. The benefits of this invention include that the polypropylene (co)polymers and blends 20 containing same provide a wider temperature processing window than conventional isotactic polypropylene. The modified polypropylene (co)polymers may also be used in large part thermoforming such as in the production of refrigerator liners and the like where conventional isotactic polypropylene is unsuitable.

- 25 The modified polypropylene (co)polymers produced in accordance with the present invention are suitable for blow moulding and we have found that they can be more readily blow moulded into containers. Furthermore, the increased melt strength makes it possible to produce large blow moulded parts through the use of the high melt strength modified polypropylene (co)polymer. Thus components currently made by rotational moulding may

now be produced by blow moulding using the modified polypropylene (co)polymer of the present invention.

Profile extrusion for example tube or pipe extrusion, using the modified polypropylene 5 (co)polymer has been found to produce a more consistent product than conventional isotactic polypropylene.

Blown films made of polypropylene are generally blown downwards using relatively expensive equipment. The modified polypropylene (co)polymers of the present invention 10 have sufficient melt strength for them to be able them to be blown upwardly using conventional polyethylene type film blowing equipment which is less expensive and generally more convenient to operate. Advantageously the modified polypropylene (co)polymers of the present invention may be used in the production of blown films.

15 The modified polypropylene (co)polymers of the present invention may also be foamed with a wider processing window than for conventional polypropylene. Either a physical or chemical blowing agent may be used. It is preferred to use carbon dioxide as a physical blowing agent to produce foams having a fine closed cell structure. Foamed pellets may be subsequently moulded to form components for use in a variety of applications such as 20 automotive door trims, rooflinings, dash boards, bumpers and the like. Applications such as in foamed packaging are also possible, including thermoformed containers, insulating cups and the like.

Waste polypropylene or waste streams containing a significant proportion of polypropylene 25 are presently difficult to recycle as conventionally a high degree of chain scission results from the recycling process. The process of the present invention may be used to upgrade recycled streams containing polypropylene by increasing the overall mechanical properties of the recycled polypropylene by the addition of initiator and monomer in accordance with the present invention.

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The present invention will now be described with reference to the following non-limiting examples. Described hereunder are the measurement techniques used in the examples and a full description of the process conditions employed. Comparative Examples are labelled CE-n.

5

Melt Strength Measurement

- Melt strengths were measured on a "Rheotens" Melt Strength Tester, Type 010.1, supplied by Gottfert Werkstoff-Prufmaschinen GmbH of Buchen, Germany. This test involves drawing an extruded strand of polymer vertically into the nip between two counter-rotating nip rollers. The strand was extruded using a Brabender Plasticord single screw extruder of screw diameter 19mm and length to diameter ratio (L/D) of 25. The extrudate exited via a right angle capillary die (2mm diameter). The temperature profile used was uniform along the length of the barrel of the extruder and the die and was set at 190°C. The nip rollers are mounted on a balance arm which allows the force in the drawing strand to be measured. The velocity of the nip rolls is increased at a uniform acceleration rate. As the test proceeds, the force increases until eventually the strand breaks. The force at breakage is termed the "melt strength".
- While there is no internationally-established standard set of test requirements for melt strength testing, comparative melt strength values obtained under the given set of test conditions provide a quantitative determination of the increase in melt strength used in the patent. The test conditions used were: die temperature 190°C, extruder output rate ~4 g/min, acceleration rate 1.2 cm/sec², draw distance 210 mm, matt finish steel rollers.

25

Dow Rheology Index

The Dow Rheology Index (DRI) is believed in the art to be a measure of the long chain branching in a polymer. It is expressed as the deviation of a viscosity parameter obtained

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from shear rheology measurements on a "branched" polymer compared with that for a linear polymer. The branched polymers have lower values of the viscosity parameter than the linear polymers (for a given relaxation parameter). The parameters are obtained by fitting the Cross model to the shear viscosity flow curves. The DRI method has been described by Lai,
5 Plumley, Butler, Knight and Kao in a paper in SPE ANTEC '94 Conference Proceedings (pp1814-1818) - "Dow Rheology Index (DRI) for Insite Technology Polyolefins (ITP): Unique Structure-Processing Relationships".

Dynamic Rheology Tests

10

The dynamic rheology tests were performed on a Rheometrics Dynamic Stress Rheometer SR200. Test conditions were: parallel plates, temperature 190°C, frequency range 0.01 to 100 rad/sec, and 3-4% strain, in a nitrogen atmosphere to prevent degradation. G' is the storage modulus representing the elasticity of the polymer melt, G" is the loss modulus which
15 represents the viscous component of the deformation. The polydispersity index is 10 to power 5 divided by the crossover modulus, which is the value of G' = G" when the G' and G" curves crossover - it is believed to be a measure of MWD. The higher G', the greater elasticity in the polymer and the higher the MW.

20 MFI

Melt flow indexes (MFI) were measured at 230°C with a 2.16 kg load according to ASTM 1238.

25 Drop Times

The drop times were determined by measuring the time taken for the polypropylene strand (cut at the die face) to drop from the die of the extruder to the floor. The die of the JSW twin screw extruder was 1140 mm above the floor. The drop time test combines the effects of

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melt viscosity, extensional viscosity, chain entanglement (as shown by die swell), and elasticity (as shown by the tendency resist neck formation). Higher melt viscosity polypropylene polymers had drop times which incorporated some additional effect due to prolonged cooling of the slower moving (falling) molten strand.

5

GPC

GPC molecular weights were determined using a Waters 150C high temperature GPC unit. 1,2,4-trichlorobenzene was used as the solvent, eluting through two Styragel HT6E linear 10 columns. The oven temperature was set at 140°C and the pump flow rate was 1.0ml/min.

Calibration was performed using narrow polydispersity polystyrene standards. All molecular weights quoted as polystyrene equivalents.

M_n= number average molecular weight

15 M_w= weight average molecular weight

M_z= viscosity average molecular weight

M_p= peak molecular weight

Twin Screw Extruder

20

The twin screw extruder used in the examples was a JSW TEX-30 with a 30 mm screw diameter and an overall L/D of 42 [comprising ten temperature controlled barrel sections (L/D 3.5, temperatures between 120 and 230°C as specified in Table 1), three unheated sampling/monitoring blocks (L/D 1.167) and a cooled feed block (L/D 3.5)) equipped with 25 two JSW TTF20 gravimetric feeders, one K-Tron KQx gravimetric additives feeder and a volumetric liquid addition pump (Fuji Techno Industries model HYM-03-08)]. The extruder was operated in either co-rotating (intermeshing self wiping) or counter rotating (intermeshing non-self wiping) modes with a throughput rate of between 5 and 20 kg/hr and screw speeds of between 100 and 400 rpm as specified in Table 1. The melt temperature and

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pressures were monitored at three points along the barrel and in the die.

Table 1 Operating conditions

Conditions		Screw Speed (rpm)	Feed Rate (kg/hr)	Temperature Profile (°C)
5	A	265	20	150°C, 175°C (by 10)
	B	265	20	180°C, 200°C (by 3), 220°C (by 7)
	C	150	5	120°C, 130°C (by 4), 180°C (by 6)
	D	265	20	140°C, 150°C (by 10)
	E	265	20	180°C, 200°C (by 4), 230°C, 240°C, 250°C, 260°C, 270°C, 280°C.
10	F	400	20	180°C, 220°C (by 10)
	G	265	20	80°C, 120°C, 140°C, 160°C, 170°C, 180°C, 200°C (by 5)
	H	150	5	80°C, 120°C, 140°C, 160°C, 170°C, 180°C, 190°C (by 3), 200°C (by 2)
	I	265	20	80°C, 120°C, 140°C, 160°C, 170°C, 180°C, 190°C (by 3), 200°C (by 2)
	J	250	20	150°C, 170°C (by 3), 180°C, 200°C, 220°C (by 5)

15

- The temperatures in the table refer to sections of the barrel of the extruder that are capable of independent temperature control. The first ten temperatures are barrel section temperatures and the last temperature indicates the temperature of the die.

20

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Table 2 - Die configuration

Condition	Die Description
1	Large 3 hole strand die - 6 mm holes
2	Small 3 hole strand die - 4 mm holes
5	Large 2 hole strand die - 6 mm holes
3	Single hole Brabender die- 10mm hole

Table 3 - Means of modifier addition

Condition	Die Description
10	Modifier added at block 4 in 2-butanone carrier solvent
α	Modifier added at block 4 in xylene carrier solvent
β	Modifier coated onto PP powder - pre tumble blended
γ	Modifier coated onto PP powder masterbatch

- 15 The overall extruder configuration and modifier conditions may be recited, for example, as condition: A1 δ .

Solvent Addition of Modifiers

- 20 The initiator, and monomer if present, was introduced as a solution in 2-butanone or xylene. The concentration of the initiator varied from 5.6% wt/wt to 8.5% wt/wt. The benzoyl peroxide and the di-toluol peroxides were both powders wetted with 25% (wt/wt) water. The monomer was present in an amount between 4 to 10% wt/wt solvent.

- 25 Increased levels of initiator were generally added by increasing the amount of solution added to the polymer melt. The additional peroxides (if any) were added with the initiator in the carrier solvent.

Solventless Addition of Modifiers

- 30 t-Butyl peroxybenzoate is a liquid. The solventless modification of the polymer was achieved by absorbing the initiator onto powdered polymer or blending it with powdered polymer at

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concentrations ranging from 5% wt/wt to 10% wt/wt to form a masterbatch. The masterbatch was added to the extruder in varying feed rates to alter the amount of additives. The amount of polymer feed was adjusted accordingly to give constant overall feed rate.

5 The stabilisers were also added as a masterbatch. The amount of stabiliser was generally kept constant at 0.33%wt/wt Irganox 1010 and 0.17%wt/wt Irgaphos 168 in the total composition.

The main polymer feed was added as either powder or pellets.

10 Single Screw Extruders

Killion

The Killion single screw extruder used in the examples was a segmented single screw extruder
15 of L/D=40 (11 barrel sections , 10 heated) and screw diameter of 31.75 mm.

Polypropylene powder, stabilisers (0.33%wt/wt Irganox 1010, 0.17%wt/wt Irgaphos 168 in total) and initiator were added to the feed throat of the single screw extruder via a twin screw K-Tron volumetric feeder.

20

Alternatively, the polypropylene powder and stabilisers were added via the K-Tron feeder and polypropylene powder, stabilisers and the modifiers were added as a master batch via a single screw APV Accurate volumetric feeder. The masterbatch contained 7.5%wt/wt benzoyl peroxide (prepared using a dispersion of benzoyl peroxide containing 25%wt/wt water).

25

The output of the extruder was ~ 1.5 kg/hr using a screw speed of 30rpm. The set barrel temperature was either (I) a flat 220°C with each barrel section and the die set at a temperature of 220°C or (ii) 230°C/190°C with the first six melting sections of the barrel set at 230°C and the next four metering sections of the barrel and the die set at 190°C. The melt

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temperature varied from 220 to 260°C.

Brabender

5 The Brabender single screw extruder used was a single screw extruder of L/D=25 (4 Barrel sections), compression ratio 2.5:1 and screw diameter of 19 mm. The die was a 4 mm rod die.

The screw speed of the extruder was 20 rpm. The set barrel temperature was 140°C, 170°C,
10 180°C, 180°C. Residences time: Start 3 min 40 sec; Middle 4 min 35 sec; and End 7 min
30 sec.

Polypropylene powder either as cryoground pellets or ex-reactor powder was mixed with the
modifiers and added to the feed throat, either flood feed or by a Brabender single screw
15 volumetric feeder.

The following commercial polypropylene (co)polymers were used in the examples. The
properties of the (co)polymers are shown in Table 4 below.

20 Table 4: Comparative data for a grade of high melt strength PP and conventional PP grades.

Example	Polymer	Polymer Description	MFI 2.16kg @ 230°C	Melt Strength cN
Control 1	Montell PF814	High melt strength polypropylene homopolymer	3	18
Control 2	Montell JE6100	Extrusion grade polypropylene homopolymer	3	3

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	Control 3	ICI	Injection moulding	14	1.8
		Australia	grade of		
		GYM 45	polypropylene		
			homopolymer		
	Control 4	ICI	Extrusion grade of	4	2.8
		Australia	polypropylene		
		GWM 22	homopolymer		
	Control 5	ICI	Thermoforming	0.8	6
		Australia	grade of		
		PXCA 6152	polypropylene		
			homopolymer		
	Control 6	ICI	Injection moulding	14	1.4
		Australia	grade of		
		LYM 120	propylene/ethylene		
			copolymer		
5	Control 7	Montell	Ex-reactor grade of	4.1	~3
		6501	injection moulding		
			polypropylene		
			homopolymer		
	Control 8	Montell	Extrusion grade of	~3.5	-
		KM6100	polypropylene		
			homopolymer		
	Control 9	Montell	Extrusion grade of	~3.5	-
		KMT6100	polypropylene		
			homopolymer		
	Control 10	Montell	Ex-reactor grade of	~3.5	-
		KM6100	polypropylene		
			powder homo-		
			polymer		
			unstabilized		

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* Melt strength and MFI were measured for a particular batch and we have found actual values vary up to 20% of these values.

Examples 1 to 5

5

GYM45 was modified in accordance with Table 5 below. GYM45 is a low molecular weight/higher MFI injection moulding grade of polypropylene homopolymer.

Table 5:

10	Example	Conditions	BPO	Styrene	Motor	Die	Drop	MFI	Melt Strength
			(wt%)	(wt%)	Current	Temp.	Time	2.16kg	(cN)
					(amps)	(°C)	(secs.)	@	230°C
	Control	-	-	0	-	-	-	14	1.8
	3								
	CE 1	B1α	0	0	13	231	8	12.2	1.5
	1	B1α	0.36	0	13	229	9.7	14.4	1.9
15	2	B1α	0.7	0	13	229	13.5	14.4	2.3
	3	B1α	0.95	0	13	230	17.1	12.5	3
	4	D3α	1.0	0	19	197	23	9.7	4
	5	D4α	0.34	0.45	21	179	22.9	9.1	6.9

20

Examples 6 to 18

GWM22 was modified in accordance with Table 6. GWM22 is an intermediate molecular weight/medium MFI extrusion grade of polypropylene homopolymer.

25

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Table 6:

Example	Conditions	BPO	Styrene	Motor	Die	Drop	MFI	Melt
		(wt%)	(wt%)	Current (amps)	Temp. (°C)	Time (secs.)	2.16kg @ 230°C	Strength (cN)
Control 4								
5	CE 2	B1α	0	0	16	239	11.3	5
	6	B1α	0.36	0	16	234	15.2	6.3
	7	B1α	0.75	0	17	238	21.8	5.9
	8	B1α	1	0	16	239	25.4	5
	9	B1α	1.3	0	20	236	25.3	5.6
10	10	B1α	0.12	0.16	16	237	8.0	4.15
	11	B1α	0.23	0.31	17	237	11.2	2.8
	12	B1α	0.46	0.61	21	238	14	1.11
	13	B1α	0.69	0.92	21	241	14.2	0.69
	14	B1α	1.22	1.63	21	248	-	18.6
15	15	E1α	0.33	0.44	17	281	20	3.6
	16	C2α	0.81	4.2	18	203	60	0.69
	17	E1α	0.31	0.40	20	275	23	3.1
	18	E1α	0.30	0.53	17	277	24	9.1

20

The increase in complex viscosity of examples 14, 16, 17 and 18 is shown in Figure 1. G' has been plotted against frequency in Figure 2.

25 The modified polypropylene's of examples 14, 16, 17 and 18 were tested for additional physical properties and it was found that the modified polypropylene's had:

		14	16	17	18	Control 4	
i)	Elasticity	1200	680	40	45	10	G'@ 0.01 rad/s (pa) rad/s (Pa)
ii)	1/Relaxation Time	~0.0013	0.085	15	18	23	Crossover Frequency(rad/sec)

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iii)	Polydispersity Index	222	39	4.4	4.7	3.7	M_w/M_n
iv)	Dow Rheology Index	192	86	2.0	5.6	0	LongChain Branching

5 Examples 19 to 26

PXCA6152 was modified in accordance with Table 7 below. PXCA6152 is a high molecular weight/low MFI thermoforming grade of polypropylene homopolymer.

10 Table 7:

Example	Conditions	BPO	Styrene	Motor	Die	Drop	MFI	Melt
		(wt%)	(wt%)	Current (amps)	Temp. (°C)	Time (secs.)	2.16kg @ 230°C	Strength (cN)
Control 5								
CE 3	F1α	0	0	17	251	14.6	1.1	-
19	B1α	0.34	0	22	244	25.9	1.3	7.4
15 20	B1α	0.68	0	23	250	22.8	1.1	11.1
21	B1α	0.8	0	24	246	30.5	0.8	14
22	B1α	1.04	0	24	247	25.3	0.65	17.7
23	F1α	0.31	0.41	21	256	24.4	0.42	17.5
24	F1α	0.47	0.63	21	264	25	0.31	-
20 25	F1α	0.55	0.73	23	269	-	0.40	-
26	F1α	0.71	0.95	22	259	25	0.35	21.3

The modified polypropylene of example 22 was tested for additional physical properties and
25 it was found that the modified polypropylene had:

- i) Elasticity 200 G' @0.01 rad/s (Pa)
- ii) 1/Relaxation Time 7.1 Crossover frequency (rad/sec)

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iii)	Polydispersity Index	3.9	M_w/M_n
iv)	Dow Rheology Index	10	Long Chain Branching

The DRI of the base polypropylene material, PXCA 6152 (an unbranched polypropylene) was 5 expected to be 0. The DRI of the modified polypropylene demonstrates a significant degree of long chain branching.

Examples 27 to 33

10 LYM120 was modified in accordance with Table 8 below. LYM120 is a low molecular weight/higher MFI injection moulding grade of PP copolymer.

Table 8:

Example	Conditions	BPO	Styrene	Motor	Die	Drop	MFI	Melt
		(wt%)	(wt%)	Current (amps)	Temp. (°C)	Time (secs.)	2.16kg @ 230°C	Strength (cN)
15	Control 6						12.2	1.4
	27	D2α	0.68	0	13	182	19.3	13.1
	28	A4α	1.08	0	19.5	200	31	9
	29	A2α	0.33	0.44	18	202	28	5.8
	30	D2α	0.32	0.42	23	185	46.5	3.8
20	31	A4α	0.42	0.55	19.5	204	31.1	6.5
	32	A4α	0.62	0.83	20	201	36.8	-
	33	A4β	0.34	0.45	16	199	25.1	4.3

Examples 34 to 42

Ex-reactor GYM45 powder was modified according to Table 9 below. GYM45 is a low molecular weight/higher MFI injection moulding grade of polypropylene homopolymer. The polypropylene was stabilized with Irganox 1010 (0.33 wt%) and Irgaphos (0.17 wt%). The 30 modifiers and stabilizers were added to the twin-screw extruder at the feed throat.

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Table 9:

Example	Conditions	BPO (wt%)	Styrene (wt%)	Motor Current (amps)	Die Temp. (°C)	Drop Time (secs.)	MFI 2.16kg @ 230°C	Melt Strength (cN)
	Control	-	-	-	-	-	14	1.8
	3							
5	CE 4	H3δ	0	0	7	209	21.5	11.3
	34	I3δ	0.38	0	14	209	12.1	13.6
	36	I3δ	0.75	0	15	210	15.0	11.8
	37	I3δ	1.5	0	15	214	20.6	10.3
	38	H3δ	0.75	0	6	209	35.5	17.6
10	39	H3δ	1.13	0	8	208	39.5	13.3
	40	H3δ	1.5	0	8	208	43.3	9.8
	41	I3δ	0.15	0.2	18	215	16.6	9.6
	42	I3δ	0.23	0.3	16	214	20.5	6.5
15								

Examples 43 to 49

GYM45 was modified in accordance with Table 10 below. GYM45 is a low molecular weight/higher MFI injection moulding grade of polypropylene homopolymer.

Table 10:

Example	Conditions	Initiator	Initiator (wt%)	Styrene (wt%)	Motor Current (amps)	Die Temp. (°C)	Drop Time (secs)	MFI 2.16kg @ 230°C
	CE 5	A3α	-	0	0	18	192	11.8
25	43	A3α	BPO	0.12	0.16	16	197	14
	44	A3α	BPO	0.21	0.28	17	200	18.8
	45	A3α	BPO	0.41	0.55	20	206	27.6
	46	A3α	BPO	0.62	0.83	22	208	32.2
	CE-6	A3α	DHBP	0.33	0.09	14.5	191	4.6
30	CE-7	A3α	DHBP	0.60	0.17	16	190	4
								117

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CE-8	A3α	DHBP	0.90	0.28	14.5	190	3.9	132	
CE-9	A3α	TBEH	0.33	0.34	16	191	9.9	18.3	
CE-10	A3α	TBEH	0.60	0.62	17	192	10	19.5	
CE-11	A3α	TBEH	0.90	0.93	17	193	10.6	17.2	
5	47	A3α	TBPB	0.30	0.30	15	194	7.8	58.3
	48	A3α	TBPB	0.68	0.70	17	198	14	47.3
	49	A3α	TBPB	0.89	0.91	19	199	15.6	38.5
	CE-12	A3α	DCP	0.08	0.09	14.5	192	4.3	48.5
10	CE-13	A3α	DCP	0.17	0.17	15	191	3.9	64.7
	CE-14	A3α	DCP	0.25	0.25	15	191	3.7	90.3
	CE-15	A3α	DLP	0.33	0.33	15	190	11.2	16.5
	CE-16	A3α	DLP	0.63	0.64	15	190	11.1	15.1
15	CE-17	A3α	DLP	0.92	0.93	15	190	11.1	18.0

15

Examples 50 to 54

LYM120 was modified in accordance with Table 11 below. LYM120 is a low molecular weight/higher MFI injection moulding grade of polypropylene copolymer.

20

Table 11:

Example	Conditions	BPO (wt%)	Motor Current (amps)	Die Temp (°C)	Drop Time (secs.)	MFI	Melt Strength
						2.16kg @ 230°C	(cN)
Control 6						12.2	1.4
25	CE 18	H3δ	0	7	209	18.4	11.3
	52	H3δ	0.75	8	210	50.5	6.9
	53	H3δ	1.13	8	210	47.0	6.6
	54	I3δ	1.5	14	217	18.5	3.8

30

Examples 55 to 61

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LYM120 was modified in accordance with Table 12 below. LYM120 is a low molecular weight/higher MFI injection moulding grade of polypropylene copolymer.

Table 12:

	Example	Conditions	Initiator	Initiator (wt%)	Styrene (wt%)	Motor Current (amps)	Die Temp. (°C)	Drop Time (secs)	MFI 2.16kg @ 230°C	Melt Strength (cN)
5	CE-19	A3α	BPO	0.00	0.00	16	195	9.9	12.8	1.1
	55	A3α	BPO	0.12	0.16	15	197	13.9	10.9	
	56	A3α	BPO	0.21	0.28	17.5	201	17.7	7.50	
	57	A3α	BPO	0.41	0.55	20	208	25.8	4.4	11.5
10	58	A3α	BPO	0.62	0.83	21	209	26.5	2.9	
	CE-20	A3α	DHB	0.08	0.09	13.5	191	4.9	52	
			P							
	CE-21	A3α	DHB	0.16	0.17	14	190	5.3	79	
			P							
	CE-22	A3α	DHB	0.28	0.30	14.5	190	5.6	114	
			P							
	CE-23	A3α	TBEH	0.31	0.32	14	192	8.6	17.8	
15	CE-24	A3α	TBEH	0.62	0.64	14	192	9	17.4	
	CE-25	A3α	TBEH	0.98	1.01	14	192	9.6	15.4	
	59	A3α	TBPB	0.30	0.31	14	196	4.6	33.8	
	60	A3α	TBPB	0.61	0.62	17	200	14.6	32.9	
	61	A3α	TBPB	0.93	0.95	17	202	15.6	23.1	
20	CE-26	A3α	DCP	0.08	0.09	13	192	5	38.6	
	CE-27	A3α	DCP	0.17	0.17	13.5	190	5.5	57.6	
	CE-28	A3α	DCP	0.27	0.28	14	190	6.2	65.9	
	CE-29	A3α	DLP	0.31	0.31	15	191	9.9	15.7	
	CE-30	A3α	DLP	0.64	0.65	13.5	190	9.8	14.8	
25	CE-31	A3α	DLP	1.00	1.01	13	190		14.8	

Examples 62 to 73

LYM120 was modified in accordance with Table 13 below. LYM120 is a low molecular weight/higher MFI injection moulding grade of polypropylene copolymer.

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Table 13:

	Example	Conditions	Initiator	Initiator (wt%)	Styrene (wt%)	Motor Current (amps)	Die Temp. (°C)	Drop Time (secs)	MFI 2.16kg @	Melt Strength (cN)
230°C										
	CE-32	A3δ	-	0		14.5	200	10	12.4	1.1
5	62	A3δ	BPO	0.11	0	14	199	10	11.5	1.1
	63	A3δ	BPO	0.23	0	16.5	201	15.1	9.0	1.2
	64	A3δ	BPO	0.45	0	16.5	203	20.6	6.6	1.7
	65	A3δ	BPO	0.68	0	17.5	206	20.6	5.4	4.1
	66	A3δ	BPO	1.13	0	18	206	18.6	5.7	3.9
10	CE-33	A3δ	DLP	0.31	0	13.5	198	9.3	13.6	-
	CE-34	A3δ	DLP	0.59	0	14	199	8.8	14.4	-
	CE-35	A3δ	DLP	0.89	0	13	196	8.9	15.0	-
	CE-36	A3δ	TBPPB	0.07	0	12	189	5.1	28.7	-
	CE-37	A3δ	TBPPB	0.15	0	12	190	5.3	31.0	-
15	CE-38	A3δ	TBPPB	0.29	0	10.5	185	6.3	92.0	0.5
	CE-39	A3δ	TBPPB	0.59	0	11	186	11.2	102.0	-
	67	A3δ	BPO	0.11	0.15	17.5	205	19.1	5.7	2.4
	68	A3δ	BPO	0.23	0.30	19.5	210	25	4.3	6.3
	69	A3δ	BPO	0.45	0.6	21.5	209	27.8	2.1	10.4
20	70	A3δ	BPO	0.90	1.2	23.5	210	26.8	1.3	12.2
	CE-40	A3δ	DLP	0.30	0.3	14.5	199	9.4	14.1	-
	CE-41	A3δ	DLP	0.59	0.6	13	197	9.3	17.2	-
	CE-42	A3δ	DLP	0.89	0.9	13	197	9.8	16.1	-
	71	A3δ	TBPPB	0.29	0.3	13.5	196	11.4	20.4	3.7
25	72	A3δ	TBPPB	0.59	0.6	16.5	200	20.2	12.9	-
	73	A3δ	TBPPB	1.18	1.2	17	202	18.8	13.9	4.5

Examples 74 to 77

30

GYM45 was modified in accordance with Table 13 below. GYM45 is a low molecular weight/higher MFI injection moulding grade of polypropylene homopolymer.

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Table 13

	Example	Conditions	Initiator	Initiator (wt %)	Motor Current (amps)	Die Temp. (°C)	Drop Time (secs)	MFI 2.16kg @ 230°C
	CE 43	A3 α	-	0	18	192	11.8	12.8
5	74	A3 α	BPO	0.23	16.5	196	10.35	17.0
	75	A3 α	BPO	0.45	17	199	11.8	17.3
	76	A3 α	BPO	0.73	17	200	15.9	15.4
	77	A3 α	BPO	0.96	18	202	17.3	14.9
	CE-44	A3 α	DHBP	0.08	15	191	3.6	96
10	CE-45	A3 α	DHBP	0.17	14.5	190	3.1	169
	CE-46	A3 α	DHBP	0.29	13	188	2.3	>100
	CE-47	A3 α	DHBP	0.30	13	188	2.2	>200
	CE-48	A3 α	DHBP	0.50	13.5	186	1.9	>200
	CE-49	A3 α	DHBP	0.57	12	186	1.9	>100
15	CE-50	A3 α	TBEH	0.30	17	197	8.8	12.0
	CE-51	A3 α	TBEH	0.64	14	188	7.2	24.4
	CE-52	A3 α	TBEH	0.98	14	189	6.8	25.6
	CE-53	A3 α	TBPB	0.31	13	186	2.6	95
	CE-54	A3 α	TBPB	0.64	13	184	2.1	238
20	CE-55	A3 α	TBPB	1.03	12	184	1.9	>250
	CE-56	A3 α	DCP	0.08	16	192	3.8	47.1
	CE-57	A3 α	DCP	0.17	14	189	2.9	121.3
	CE-58	A3 α	DLP	0.32	16	191	10.9	14.5
	CE-59	A3 α	DLP	0.64	16	190	10.6	17.9
25	CE-60	A3 α	DLP	0.95	16	189	10.4	16.6

Examples 78 to 82

- 30 LYM120 was modified in accordance with Table 14 below. LYM120 is a low molecular weight/higher MFI injection moulding grade of polypropylene copolymer.

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Table 14:

Example	Conditions	Initiator	Initiator (wt%)	Motor Current (amps)	Die Temp. (°C)	Drop Time (secs)	MFI	
							2.16kg @ 230°C	
	Control 24	A3α	BPO	0	16	195	9.9	12.8
5	78	A3α	BPO	0.25	15	196	10.3	13.4
	79	A3α	BPO	0.47	14	198	13.2	13.1
	80	A3α	BPO	0.64	16	198	13.9	12.9
	81	A3α	BPO	0.70	17	201	13.6	11.9
	82	A3α	BPO	0.94	18	198	13.5	10.6
10	CE-61	A3α	DHBP	0.09	14	190	3.6	80
	CE-62	A3α	DHBP	0.16	13	188	3.1	160
	CE-63	A3α	DHBP	0.26	13	187	2.8	250
	CE-64	A3α	TBEH	0.29	15	193	7	
	CE-65	A3α	TBEH	0.60	13	192	6.7	19.4
15	CE-66	A3α	TBEH	1.02	12	191	6	22.6
	CE-67	A3α	TBPB	0.30	13	186	3.6	85
	CE-68	A3α	TBPB	0.61	12	184	3.8	173
	CE-69	A3α	TBPB	0.92	12	184	3.4	>250
	CE-70	A3α	DCP	0.08	16	192	4.5	14.6
20	CE-71	A3α	DCP	0.17	13	190	3	107
	CE-72	A3α	DCP	0.25	13	188	3	131
	CE-73	A3α	DLP	0.32	14.5	191	9.5	14.3
	CE-74	A3α	DLP	0.68	15	191	9.4	15.8
	CE-75	A3α	DLP	0.98	14.5	193	8.7	21.0

25

Examples 83 to 92

LYM120 was modified in accordance with Table 15 below. LYM120 is a low molecular weight/higher MFI injection moulding grade of polypropylene copolymer.

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Table 15:

Example	Conditions	Initiator	Initiator (wt%)	Styrene (wt%)	Mole Ratio Sty/Init	Motor Current (amps)	Die Temp. (°C)	Drop Time (secs)	MFI	Melt
									2.16kg @ 230°C	(cN)
CE-76	A3α	0	0	0	-	16	195	9.9	12.8	1.1
5	83	A3α	BPO	0.43	0.19	1.04	16	207	18.5	6.9
	84	A3α	BPO	0.41	0.37	2.07	19	207	22.5	4.9
	85	A3α	BPO	0.41	0.55	3.11	20	208	25.8	4.4
	86	A3α	BPO	0.43	0.76	4.14	21	210	25.4	3.9
	87	A3α	BPO	0.45	0.99	5.18	20	205	28.3	4.4
10	88	A3α	TBPB	0.56	0.19	0.78	13	192	8.9	71
	89	A3α	TBPB	0.55	0.37	1.57	14	196	11.0	38
	90	A3α	TBPB	0.61	0.62	1.91	17	200	14.6	33
	91	A3α	TBPB	0.54	0.73	3.14	16	201	16.0	19.2
	92	A3α	TBPB	0.58	0.97	3.92	19	204	16.2	15.7
CE-77	A3α	DHB	0.27	0.1	0.84	13	190	4.4	187	-
15	CE-78	A3α	DHB	0.25	0.18	1.69	14	191	5.3	125
	CE-79	A3α	DHB	0.28	0.30	3.04	14.5	190	5.6	114
	CE-80	A3α	DHB	0.27	0.40	3.38	15	193	6.2	116
	CE-81	A3α	DHB	0.28	0.50	4.22	14	192	6.1	118
			P							

20

Examples 93 to 97

GYM45 was modified in accordance with Table 16 below. GYM45 is a low molecular weight/higher MFI injection moulding grade of polypropylene homopolymer.

25

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Table 16:

Example	Conditions	Initiator	Initiator (wt%)	Styrene (wt%)	Mole Ratio	Motor Current Sty/Init (amps)	Die Temp (°C)	Drop Time (secs)	MFI	
									2.16kg @ 230°C	
5	CE-82	A3α	-	0.00	0.00	0.00	18	192	11.8	12.75
	93	A3α	BPO	0.36	0.16	1.04	18	202	18.1	11.17
	94	A3α	BPO	0.41	0.37	2.07	18	209	23.5	6.38
	95	A3α	BPO	0.41	0.55	3.11	20	206	27.6	5.62
	96	A3α	BPO	0.43	0.76	4.14	22	209	25.7	4.05
	97	A3α	BPO	0.40	0.89	5.18	21	207	31.2	4.27

10

Example 98 to 105

LYM120 was modified in accordance with Table 17 below. LYM120 is a low molecular weight/higher MFI injection moulding grade of polypropylene copolymer.

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Table 17:

Example	Conditions	Initiator #1 wt%	Initiator #2 wt%	Mole ratio	Monomer Init#1/Init#2	Mole ratio	Monomer /tot init	Motor Current (amps)	Drop time (secs)	Die temp. °C	MFI mp.	Melt strength (cN)
5 Initiator#1=BPO, Initiator #2=DHBP, Monomer=Styrene												
98	A3δ	0.43	0.06	9.08	0.57	2.81	19	17	204	6.9	3.8	
99	A3δ	0.43	0.11	4.54	0.58	2.57	19	14.8	203	10.5	2.9	
100	A3δ	0.43	0.17	3.03	0.58	2.37	18	14	201	16.9	-	
Initiator #1=BPO, Initiator#2=TBPB, Monomer=Styrene												
10	101	A3δ	0.43	0.11	3.07	0.58	2.36	20	18.7	208	5	5.5
102	A3δ	0.43	0.22	1.53	0.58	1.92	17	14.2	204	9.3	4.2	
103	A3δ	0.43	0.34	1.02	0.59	1.62	18	14	204	11.6	-	
104	A3δ	0.43	0.45	0.77	0.60	1.40	18	11.8	201	19.5	-	
105	A3δ	0.43	0.34	1.02	0.94	2.59	21	17	207	7.5	6.3	

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Example 108 and 109

Montell 6501 was modified in accordance with Table 19 below on the Killion screw extruder described above.

5

Sample	Barrel Temp (°C)	Extruder Output (kg/hr)	BPO wt%	Styrene wt%	Motor Current (amps)	Die Temp °C)	Drop Time (secs)	MFI (g/10 min)
Control	-	-	-	-	-	-	-	4.1
7								
CE 84	220	1.4	0	0	6	256	17	4.1
	flat							
10 108	220	1.4	2.1	0.25	6	260	35	2.2
	flat							
109	220	1.4	4.2	0.5	7.5	263	33	0.40
	flat							

Examples 40, 41, 7, 12, 28, 29, 31 and 14

15

GPC molecular weights were determined using a Waters 150C high temperature GPC unit. 1,2,4-trichlorobenzene was used as the solvent, eluting through two Ultrastyragel linear columns. The oven temperature was set at 140°C and the pump[flow rate was 1.0 ml/min.

20 Calibration was performed using narrow polydispersity polystyrene standards. All molecular weights quoted as linear polystyrene equivalents.

Mn=number average molecular weight

Mw=weight average molecular weight

25 Mz=viscosity average molecular weight

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M_p=peak molecular weight

Error are quoted as two times the standard deviation between duplicate injections.

5 Table 20:

Exam ple	Cond. No.	BPO (wt%)	Sty (wt%)	MFI (g/10 min)	Melt (cN)	Mn l) x 10 ⁻³	Mw l) x 10 ⁻³	Mz l) x 10 ⁻³	M _p l) x 10 ⁻³
Intermediate Molecular Weight PP Homopolymer (GWM 22)									
10	Contr ol	-	-	-	4.5	2.8	55	295	1200
	4								
	8	B1 α	1.0		5.0	6.9	80	425	1415
	10	B1 α	0.12	0.16	4.15	-	90	405	1200
15	11	B1 α	0.23	0.31	2.80	5.0	75	415	1400
	12	B1 α	0.46	0.61	1.11	-	70	555	2200
	13	B1 α	0.69	0.92	0.69	18.6	85	575	2200
	110	C2 α	1.50		3.8	-	75	430	1560
	111	C2 α	2.23		3.2	-	75	430	1700
20	112	C1 α	0.37	0.49	2.22	-	85	565	2035
	113	C1 α	0.60	0.80	1.00	19.4	85	690	2575
	114	C1 α	0.32	1.65	4.50	-	80	505	1835
	115	C1 α	0.47	2.45	1.58	-	90	605	2160
	116	C1 α	0.81	4.19	0.69	18.8	85	675	2610
25	Low Molecular Weight PP Copolymer (PXCA 6152)								
	Contr ol	-	-	-	12.4	1.4	45	230	720

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5										
22	B1 α	1.04		0.65	17.7	110	485	1615	195	
27	A2 α	0.33	0.44	-	7.4	65	325	1045	140	
61	A2 α	0.41	0.55	4.4	11.5	60	325	1315	125	
5	62	A2 α	0.62	0.83	2.9	-	70	460	2435	135
-	D2 α	0.28	0.38	-	-	80	555	2875	160	
28	D2 α	0.32	0.44	-	9.0	120	640	4130	140	
78	A3 α	0.25		13.4	-	70	315	1330	135	
79	A3 α	0.47		13.1	-	65	320	1380	130	
10	80	A3 α	0.64		12.9	-	65	360	1975	130
27	D3 α	0.68		13.1	2.3	70	445	1865	140	

*Errors in the molecular weight are generally less than 30% of the quoted value, as is usual in high temperature GPC under the conditions employed.

15

Examples 110 to 114

GWM22 and KM6100 were modified in accordance with Table 20 below.

Table 20: Effect of feed throat addition of BPO on the modification of prestabilised PP
 20 homopolymer [KM6100 or GWM22] *a*

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Control								~3.5	2.5
8									
CE-85	J3δ	0	0	27	11	240	3.7	2.5	
112	J3δ	2.0	0.41	26	15	235	4.9	3.9	
5	113	J3δ	3.9	0.81	27	19	241	3.3	6.7
	114	J3δ	6.2	1.28	28	19	241	2.7	9.5

a: BPO added to pellet feed in PP powder derived from cryoground prestabilised PP pellets

10

Examples 115 to 117

KMT6100 was modified in accordance with Table 22 below. KMT6100 is a prestabilised PP 15 copolymer.

Table 22: Effect of feed throat addition of BPO on the modification of prestabilised PP copolymer [KMT6100]

a: BPO added to pellet feed in PP powder derived from cryoground prestabilised PP pellets

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Examples 118 to 121

KM6100u was modified with para-toluoyl peroxide (PTP) and BPO in accordance with Table 23 below. The KM6100u was stabilized with Irganox 1010 (0.33wt%) and Irgaphos 168 5 (0.17 wt%) which were added to the main feed throat of the extruder.

Table 23:

Example	Conditions	Peroxide	Peroxide (wt%)	Motor Current (amps)	Drop Time (sec)	Die Temp °C	MFI (g/10 min.)	Melt Str. (cN)
	Control						~3.5	~2.5
10	10							
	CE-87	J3δ	-	0	22	9	240	5.2
	118	J3δ	BPO	1.0	22	17	252	5.2
	119	J3δ	PTP	1.0	21	16	240	5.2
	120	J3δ	PTP	1.5	22	18	239	3.9
15	121	J3δ	PTP	2.0		24	243	3.4
								14.0

Examples 122 to 128

PXCA6152 was modified with mixed initiator systems in accordance with Table 24 below.

5 Table 24:

Example	Conditions	Init #1	Init # 2	Mole ratio	Motor Current Init#1/I _n (amps)	Drop Time (sec)	Die Temp °C	MFI (g/10 min.)	Melt Str. (cN)
Control									
5								0.8	6
10 CE-88	B3α	0	0	-	-	22	14	255	0.9
122	B3α	0.87	0	-	-	24	20	257	1.28
									5.1
									14.2
15	123	B3α	0.87	0.045	23.2	22	19	251	4.0
	124	B3α	0.87	0.064	16.3	20	16	250	5.4

Initiator #1 = BPO, Initiator # 2 = DHBP

125 B3 α 0.87 0.084 12.4 20 15 248 6.4 0.2

Initiator #1 = BPO, Initiator # 2 = TBPPB

5	126	B3 α	0.87	0.006	129.4	22	16	254	3.4	-
	127	B3 α	0.87	0.012	64.7	20	16	249	3.9	0.5
	128	B3 α	0.87	0.019	40.9	20	16	249	6.4	0.1

10

Examples 129 to 132

15 Cryoground PXCA6152 in the form in the form of a powder was modified with mixed initiator systems according to Table 25 below.

Table 25: Effect of Mixed Initiators on the Modification of PXCA6152 Powder α (cryoground pellets)

Example	Conditions	Init #1	Init # 2	Mole ratio	Motor	Drop	Die	MFI	Melt Str.
	wt%	wt%	Init#1/Init #2		Current (amps)	Time (sec)	Temp °C	(g/10 min.)	(cN)
Control									
5	CE-89	B3δ	0	0	-	25	13	256	1.0
5	Initiator #1 = BPO, Initiator # 2 = DHBP					23	17	253	1.8
129	B3δ	0.86	0	-					10.8
10	130	B3δ	0.87	0.004		23	17	254	2.1
									8.8
Initiator #1 = BPO, Initiator # 2 = TBPPB									
131	B3δ	0.87	0.016			24	18	253	2.6
15	132	B3δ	0.87	0.026		23	17	253	3.6
									8.2
									7.7

Examples 79 and 85

20 LYM120 was modified in accordance with Table 26 below.

Table 26:

Example	Conditions	BPO wt%	Monomer/ Coagent	Monomer/ er wt%	Motor Current (amps)	Drop Time (sec)	Die Temp °C	MFI (g/10 min.)	Melt Str. (cN)
Control								12.2	1.4
6									
5	CE-76	A α	0	none	0	16	10	195	12.8
79	A α	0.47	none	0	0	14	13	198	13.2
85	A α	0.41	Styrene	0.54	20	25.8	208	4.4	11.6

10 Examples 133

GYM 22 was modified in accordance with Table 27 below.

Table 27:

Example	Conditions	BPO	Monomer/ Coagent	Monomer wt%	Motor Current (amps)	Drop Time (sec)	Die Temp °C	MFI (g/10 min.)	cN	Melt Str.
Control										
5	4							4.5	2.8	
CE-2	B1α	0	none	-	16	11	239	5	-	
6	B1α	0.36	none	-	16	15	234	6.3	3	
133	B1α	0.34	Styrene	0.45	21	29	250	1.72	-	

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Examples 134 to 137

Cryoground KM6100 in the form of a powder was modified on a Brabender single screw extruder in accordance with the general description of the Brabender SSE above and Table 5 28 below. The initiator was added at the feed throat of the SSE along with the stabilizers (0.33 wt% Irganox 1010 and 0.17 wt% Irgaphos 168).

Table 28:

10	Example	Peroxide	Peroxide	MFI
		Type	(wt%)	(g/10 min.)
	Control 8	-	0	3.5
	134	BPO	1	2.9
	135	PTP	0.5	3.8
	136	PTP	1	3.3
15	137	PTP	2	2.0

* PTP- Paratoluol Peroxide (bis paramethyl benzoyl peroxide)

20 Example 138

PXCA6152 was modified in accordance with Table 29 below.

Table 29: Modification of PXCA6152 Pellets

Example	Conditions	BPO	Styrene	Motor	Drop	Die	MFI	Melt
		wt%	wt%	Current	Time	Temp	(g/10 min.)	Str.
				(amps)	(sec)	°C		(cN)
25	Control						0.8	6.0
	5							
CE-94	B3α	0	-	22	14	255	0.9	5.1
138	B3α	0.51	0.68	24	19	279	0.6	21.0

Examples 139 to 143

The modified PXCA6152 produced according to Example 138 was melt mixed with GYM45 in accordance with Table 30.

Table 30: Blends of the Modified PP with other PP Homopolymers

Example	Conditions	PP#1 (wt%)	PP#1 (wt%)	Motor Current (amps)	Drop time (seconds)	Die Temp °C	MFI	Melt Strength
5	CE-95 A3	Control 5 (5)	Control 3 (95)	18	11	206	9.9	
		Control 5 (10)	Control 3 (90)	19	12	202	7.8	
		Control 5 (15)	Control 3 (95)	19	13	202	7.0	
		Control 5 (20)	Control 3 (80)	19	14	202	5.5	
		Control 5 (25)	Control 3 (75)	19	15	202	4.7	2.6
		138 (5)	Control 3 (95)	19	15	199	9.3	
10	139 A3	138 (10)	Control 3 (90)	19	16	201	7.4	
		138 (15)	Control 3 (85)	19	18	203	6.6	
		138 (20)	Control 3 (80)	20	19	204	5.0	
		138 (25)	Control 3 (75)	19	20	207	4.7	5.2
		A3	Control 4 (100)	20	15	208	5.3	
		A3	Control 4 (95)	21	18	207	3.7	
15	CE-100 A3	138 (5)	Control 4 (90)	21	19	206	3.5	
		138 (10)	Control 4 (85)	22	21	210	2.3	
		138 (15)	Control 4 (80)	22	21	212	2.6	
		138 (20)	Control 4 (75)	23	23	213	2.4	7.7
		A3	Control 4 (75)					
20	148							

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Examples of Carbon Dioxide Foaming of Modified PP

The equipment used for foaming the polypropylene (from earlier examples) was a tandem extrusion line made up of an Leitritz twin screw extruder (34 mm screw diameter, co-5 rotating, with 11 barrel sections) connected via a melt pipe to a single screw extruder (43 mm screw diameter). CO₂ was introduced into barrel six of the twin screw extruder. The gassed polymer was then cooled slowly in the single screw extruder.

Example	MFI (g/10 min)	Melt Strength (cN)	Foaming Temp (°C)	Av Foam Density (g/cc)	Av Cell Size (μm)
10 Control 1	3	18	166 to 159	0.058	550
25	0.4	-	169 to 159	0.044	300
31	6.5	11.2	167 to 161	0.051	280

Non high melt strength grades of polypropylene have foam temperature processing windows 15 of less than 1°C.

Foamed examples 13 and 17 both has a fine closed cell structure.

Examples of thermoforming

20

The modified polypropylene produced in Example 69 was extruded on a Welex single screw extruder through a sheet die to produce a sheet 78cm wide and ~1.25 mm thick. The sheet was fed to a Gabler F702 continuous thermoformer to produce margarine tubs. Tubs produced from the modified PP sample had a crush strength of 25 kg after 1 hour. No 25 appreciable sag was noticed of the PP sheet during the process.

Blow Moulding

The modified polypropylene of Example 5 was blow moulded on Bekum blow moulder fitted with a general purpose polyolefin screw using a 750 ml screw top bottle mould, (radially non symmetrical bottle with waist). The mould temperature was 0°C.

5

The blow mouldability of the modified injection moulding grade of PP was compared against that of a commercial low melt flow index PP homopolymer (ICI GWM110 of MFI = 1.5).

It was found that the modified PP homopolymer (MFI = 9.1 and Melt Strength = 6.9 cN) 10 could be easily blow moulded into 750 ml bottles. Conventional PP of similar MFI could not be successfully blow moulded. The modified PP gave similar performance to an extrusion grade PP of low MFI.

The results are very promising where a higher MFI PP could be used to blow bottles. This 15 possibly opens up the opportunity to produce large blow moulded parts through use of a high melt strength modified PP which has been tailored to have an MFI acceptable to blow moulding (ie 1-2 MFI)

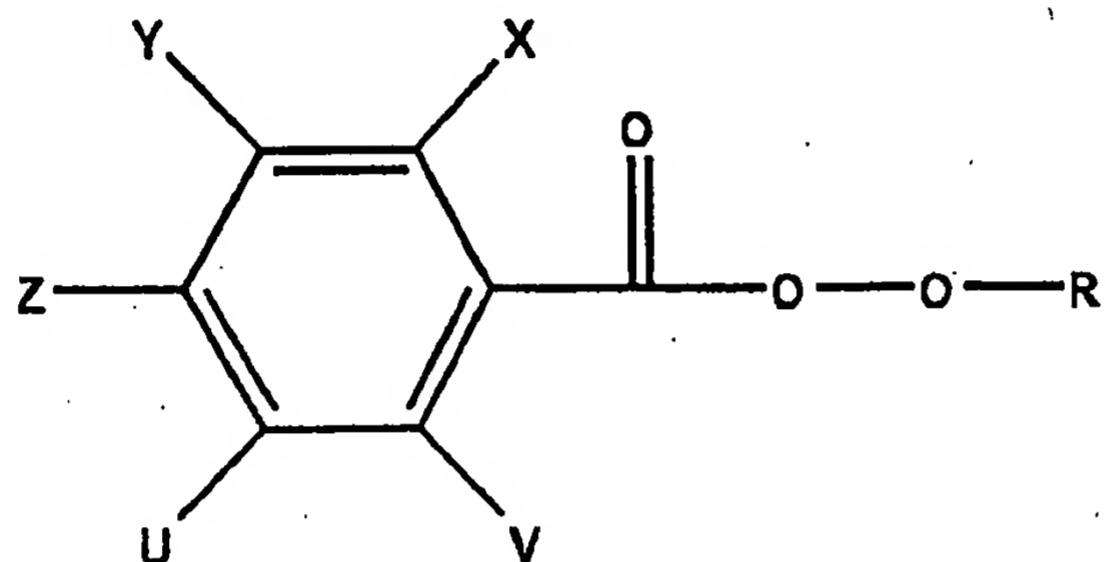
Those skilled in the art will appreciate that the invention described herein is susceptible to 20 variations and modifications other than those specifically described. It is to be understood that the invention includes all such variations and modifications which fall within its spirit and scope. The invention also includes all of the steps, features, compositions and compounds referred to or indicated in this specification, individually or collectively, and any and all combinations of any two or more of said steps or features.

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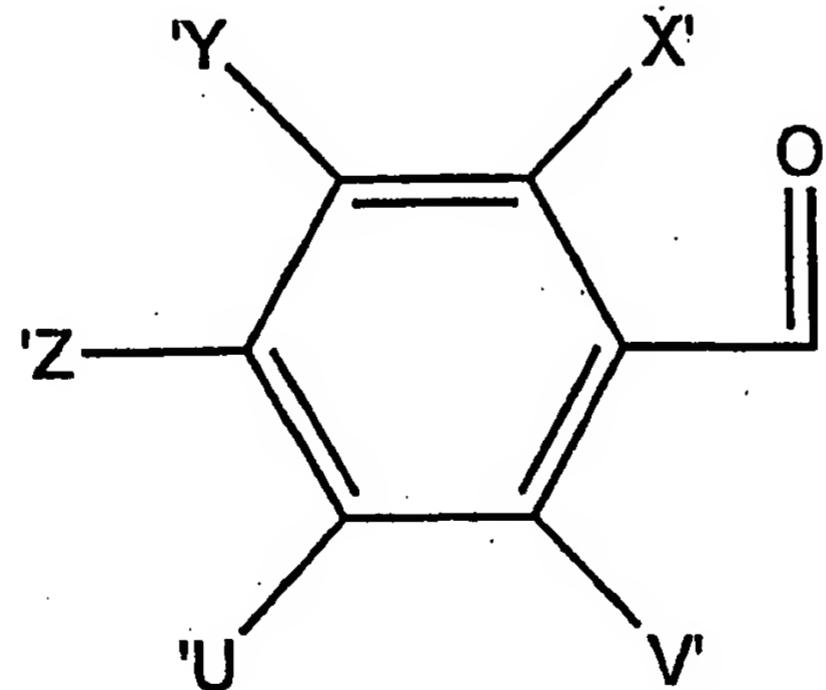
CLAIMS

1. A process for modifying a polypropylene (co)polymer wherein said process comprises melt mixing the polypropylene (co)polymer in the presence of an initiator wherein said initiator is selected from the group defined by formula 1.



Formula 1

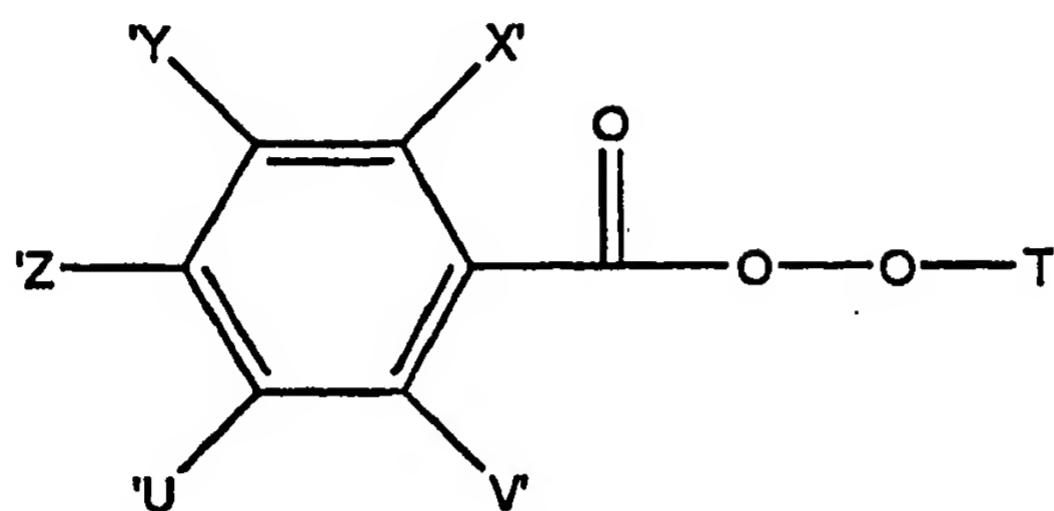
wherein R is selected from the group consisting of optionally substituted C₁ to C₁₈ acyl,
10 optionally substituted C₁ to C₁₈ alkyl, aroyl defined by formula 2,



Formula 2

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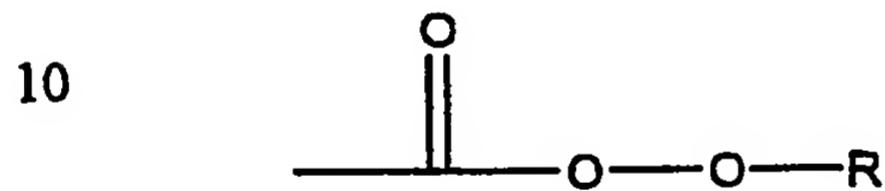
and compounds of formula 3,



Formula 3

5

wherein U, V, X, Y, Z, U', V', X', Y' and Z' are independently selected from the group consisting hydrogen; halogen; C1-C18 alkyl; C1-C18 alkoxy, aryloxy, acyl, acyloxy, aryl, carboxy, alkoxycarbonyl, aryloxycarbonyl, trialkyl silyl, hydroxy, or a moiety of formula 4,

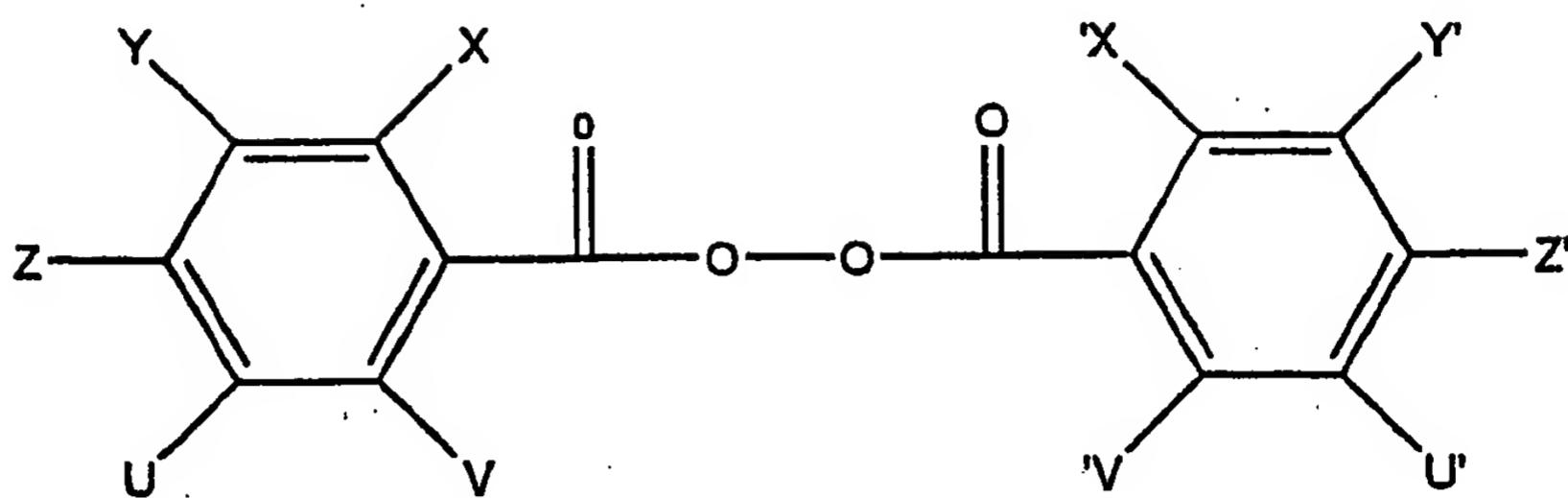


Formula 4

and wherein T is alkylene.

2. A process according to claim 1 wherein the initiator is selected from compounds of
15 formula 6.

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**Formula 6**

5 where X, Y, Z, U, V, X', Y', Z', U', V' are independently selected from the group consisting of hydrogen and C₁ - C₁₈ alkyl where at least one of X, Y, Z, U, V and X', Y', Z', U', V' are not hydrogen.

3. A process according to claim 2 wherein the initiator is selected from the group
 10 consisting of Dibenzoyl peroxide, o,o'-Bis(methylbenzoyl) peroxide, p,p'-Bis(methylbenzoyl) peroxide, M,M'-Bis(methylbenzoyl) peroxide, o,m'-Bis(methylbenzoyl) peroxide, o,p'-Bis(methylbenzoyl) peroxide, m,p'-Bis(methylbenzoyl) peroxide, Bis(ethylbenzoyl) peroxide (all isomers), Bis(propylbenzoyl) peroxide (all isomers), Bis(butylbenzoyl) peroxide (all isomers), Bis(pentylbenzoyl) peroxide (all isomers), Bis(hexylbenzoyl) peroxide (all isomers),
 15 Bis(heptylbenzoyl) peroxide (all isomers), Bis(octylbenzoyl) peroxide (all isomers), Bis(nonylbenzoyl) peroxide (all isomers), Bis(methoxybenzoyl) peroxide (all isomers), Bis(ethoxybenzoyl) peroxide (all isomers), Bis(propoxybenzoyl) peroxide (all isomers), Bis(butoxybenzoyl) peroxide (all isomers), Bis(pentoxybenzoyl) peroxide (all isomers), Bis(hexyloxybenzoyl) peroxide (all isomers), Bis(heptyloxybenzoyl) peroxide (all isomers),
 20 Bis(octyloxybenzoyl) peroxide (all isomers), Bis(nonyloxybenzoyl) peroxide (all isomers), Bis(chlorobenzoyl) peroxide (all isomers), Bis(fluorobenzoyl) peroxide (all isomers), Bis(bromobenzoyl) peroxide (all isomers), Bis(dimethylbenzoyl) peroxide (all isomers), Bis(trimethylbenzoyl) peroxide (all isomers), Bis(tert-butylbenzoyl)peroxide (all isomers), Bis(di-

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tert-butylbenzoyl)peroxide (all isomers), Bis(tertbutoxybenzoyl)peroxide (all isomers), Bis(ditrimethylsilylbenzoyl) peroxide (all isomers), Bis(heptafluoropropylbenzoyl) peroxide (all isomers), Bis(2,6-dimethyl-4-trimethylsilyl benzoyl) peroxide and isomers, 2,2'(dioxydicarbonyl) bis - Benzoic acid dibutyl ester where the term "all isomers" refers to any variation in the position 5 of the ring substituent as well as the structure of the substituent itself i.e. for propyl; n-propyl and isopropyl.

4. A process according to claim 1 wherein the initiator is selected from the group consisting of tert-butyl perbenzoate, tert-butyl (methyl)perbenzoate (all isomers), tert-butyl 10 (ethyl)perbenzoate (all isomers), tert-butyl (octyl)perbenzoate (all isomers), tert-butyl (nonyl)perbenzoate (all isomers), tert-amyl perbenzoate, tert-amyl (methyl)perbenzoate (all isomers), tert-amyl (ethyl)perbenzoate (all isomers), tert-amyl (octyl)perbenzoate (all isomers), tert-amyl (nonyl)perbenzoate (all isomers), tert-amyl (methoxy)perbenzoate (all isomers), tert-amyl (octyloxy)perbenzoate (all isomers), tert-amyl (nonyloxy)perbenzoate (all isomers), 2- 15 ethylhexyl perbenzoate, 2-ethylhexyl (methyl)perbenzoate (all isomers), , 2-ethylhexyl (ethyl)perbenzoate (all isomers), 2-ethylhexyl (octyl)perbenzoate (all isomers), , 2-ethylhexyl (nonyl)perbenzoate (all isomers), 2-ethylhexyl (methoxy)perbenzoate (all isomers), 2-ethylhexyl (ethoxy)perbenzoate (all isomers), 2-ethylhexyl (octyloxy)perbenzoate (all isomers), 2- 20 ethylhexyl (nonyloxy)perbenzoate (all isomers).

20

5. A process according to claim 1 wherein the initiator is selected from the group consisting of Bis (tertbutylmonoperoxy phthaloyl) diperoxy terephthalate, Bis (tertamylmonoperoxy phthaloyl) diperoxy terephthalate diacetyl phthaloyl diperoxide, dibenzoyl phthaloyl diperoxide, bis(4 methylbenzoyl) phthaloyl diperoxide, diacetyl terephthaloyl di 25 peroxide, dibenzoyl terephthaloyl diperoxide, Poly[dioxycarbonyldioxy(1,1,4,4-tetramethyl-1,4-butanediyil)] peroxide.

6. A process according to claim 1 wherein the initiator has a 0.1 hour half life in the range 100 - 170°C.

30

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7. A process according to claim 1 wherein the initiator is present in the range of from 0.004 to 0.25 moles of initiator per kg of the polypropylene homopolymer or copolymer.
8. A process according to claim 1 wherein the initiator is present in the range of from 5 0.006 to 0.10 moles of initiator per kg of the polypropylene homopolymer or copolymer.
9. A process according to claim 1 wherein the initiator is present in the range of from 0.01 to 0.05 moles of initiator per kg of the polypropylene homopolymer or copolymer.
10. A process according to claim 1 wherein there is no added monomer and the initiator is selected from the group consisting of Dibenzoyl peroxide, o,o'-Bis(methylbenzoyl) peroxide, p,p'-Bis(methylbenzoyl) peroxide, o,o'-Bis(methylbenzoyl) peroxide, o,m'-Bis(methylbenzoyl) peroxide, o,p'-Bis(methylbenzoyl) peroxide, m,p'-Bis(methylbenzoyl) peroxide, Bis(ethylbenzoyl) peroxide (all isomers), Bis(propylbenzoyl) peroxide (all isomers), Bis(butylbenzoyl) peroxide (all 15 isomers), Bis(pentylbenzoyl) peroxide (all isomers), Bis(hexylbenzoyl) peroxide (all isomers), Bis(heptylbenzoyl) peroxide (all isomers), Bis(octylbenzoyl) peroxide (all isomers), Bis(nonylbenzoyl) peroxide (all isomers), Bis(methoxybenzoyl) peroxide (all isomers), Bis(ethoxybenzoyl) peroxide (all isomers), Bis(propoxybenzoyl) peroxide (all isomers), Bis(butoxybenzoyl) peroxide (all isomers), Bis(pentoxybenzoyl) peroxide (all isomers), 20 Bis(hexyloxybenzoyl) peroxide (all isomers), Bis(heptyloxybenzoyl) peroxide (all isomers), Bis(octyloxybenzoyl) peroxide (all isomers), Bis(nonyloxybenzoyl) peroxide (all isomers), Bis(chlorobenzoyl) peroxide (all isomers), Bis(fluorobenzoyl) peroxide (all isomers), Bis(bromobenzoyl) peroxide (all isomers), Bis(dimethylbenzoyl) peroxide (all isomers), Bis(trimethylbenzoyl) peroxide (all isomers), Bis(tert-butylbenzoyl)peroxide (all isomers), Bis(di-25 tert-butylbenzoyl)peroxide (all isomers), Bis(tertbutoxybenzoyl)peroxide (all isomers), Bis(dtrimethylsilylbenzoyl) peroxide (all isomers), Bis(heptafluoropropylbenzoyl) peroxide (all isomers), Bis(2,4-dimethyl-6- trimethylsilyl benzoyl) peroxide and isomers tert-amyl perbenzoate, tert-amyl (methyl)perbenzoate (all isomers), tert-amyl (ethyl)perbenzoate (all isomers), tert-amyl (octyl)perbenzoate (all isomers), tert-amyl (nonyl)perbenzoate (all isomers), tert-amyl (methoxy)perbenzoate (all isomers), tert-amyl (octyloxy)perbenzoate (all isomers), tert-amyl 30

(nonyloxy)perbenzoate (all isomers), Bis (tertamylmonoperoxy phthaloyl) diperoxy terephthalate, diacetyl phthaloyl diperoxide, dibenzoyl phthaloyl diperoxide, bis(4-methylbenzoyl) phthaloyl diperoxide, diacetyl terephthaloyl di peroxide and dibenzoyl terephthaloyl diperoxide.

- 5 11. A process according to claim 10 wherein the initiator is more preferably the initiators are selected from the group consisting of dibenzoyl peroxide, o,o'-Bis(methylbenzoyl) peroxide, p,p'-Bis(methylbenzoyl) peroxide, M,M'-Bis(methylbenzoyl) peroxide, o,m'-Bis(methylbenzoyl) peroxide, o,p'-Bis(methylbenzoyl) peroxide, m,p'-Bis(methylbenzoyl) peroxide.
- 10 12. A process according to claim 1 wherein the initiator is used in combination with a monomer.
13. A process according to claim 12 wherein the monomer is a monoene monomer.
- 15 14. A process according to claim 12 wherein the monomer is styrene.
15. A process according to claim 12 wherein the initiator is selected from the group consisting of Dibenzoyl peroxide, o,o'-Bis(methylbenzoyl) peroxide, p,p'-Bis(methylbenzoyl) peroxide, M,M'-Bis(methylbenzoyl) peroxide, o,m'-Bis(methylbenzoyl) peroxide, o,p'-Bis(methylbenzoyl) peroxide, m,p'-Bis(methylbenzoyl) peroxide, Bis(ethylbenzoyl) peroxide (all isomers), Bis(propylbenzoyl) peroxide (all isomers), Bis(butylbenzoyl) peroxide (all isomers), Bis(pentylbenzoyl) peroxide (all isomers), Bis(hexylbenzoyl) peroxide (all isomers), Bis(heptylbenzoyl) peroxide (all isomers), Bis(octylbenzoyl) peroxide (all isomers), Bis(nonylbenzoyl) peroxide (all isomers), Bis(methoxybenzoyl) peroxide (all isomers),
20 Bis(ethoxybenzoyl) peroxide (all isomers), Bis(propoxybenzoyl) peroxide (all isomers), Bis(butoxybenzoyl) peroxide (all isomers), Bis(pentoxybenzoyl) peroxide (all isomers), Bis(hexyloxybenzoyl) peroxide (all isomers), Bis(heptyloxybenzoyl) peroxide (all isomers), Bis(octyloxybenzoyl) peroxide (all isomers), Bis(nonyloxybenzoyl) peroxide (all isomers), Bis(chlorobenzoyl) peroxide (all isomers), Bis(fluorobenzoyl) peroxide (all isomers),
25 Bis(bromobenzoyl) peroxide (all isomers), Bis(dimethylbenzoyl) peroxide (all isomers),
30 Bis(dimethylbenzoyl) peroxide (all isomers), Bis(dimethylbenzoyl) peroxide (all isomers),

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Bis(trimethylbenzoyl) peroxide (all isomers), Bis(tert-butylbenzoyl)peroxide (all isomers), Bis(di-tert-butylbenzoyl)peroxide (all isomers), Bis(tert-butoxybenzoyl)peroxide (all isomers), Bis(ditrimethylsilylbenzoyl) peroxide (all isomers), Bis(heptafluoropropylbenzoyl) peroxide (all isomers), Bis(2,4-dimethyl-6- trimethylsilyl benzoyl) peroxide and isomers, 2,2'(dioxydicarbonyl)
5 bis - Benzoic acid dibutyl ester, tert-butyl perbenzoate, tert-butyl (methyl)perbenzoate (all isomers), tert-butyl (ethyl)perbenzoate (all isomers), tert-butyl (octyl)perbenzoate (all isomers), tert-butyl (nonyl)perbenzoate (all isomers), tert-amyl perbenzoate, tert-amyl (methyl)perbenzoate (all isomers), tert-amyl (ethyl)perbenzoate (all isomers), tert-amyl (octyl)perbenzoate (all isomers), tert-amyl (nonyl)perbenzoate (all isomers), tert-amyl (methoxy)perbenzoate (all
10 isomers), tert-amyl (octyloxy)perbenzoate (all isomers), tert-amyl (nonyloxy)perbenzoate (all isomers), 2-ethylhexyl perbenzoate, 2-ethylhexyl (methyl)perbenzoate (all isomers), , 2-ethylhexyl (ethyl)perbenzoate (all isomers), 2-ethylhexyl (octyl)perbenzoate (all isomers), , 2-ethylhexyl (nonyl)perbenzoate (all isomers), 2-ethylhexyl (methoxy)perbenzoate (all isomers), 2-ethylhexyl (ethoxy)perbenzoate (all isomers), 2-ethylhexyl (octyloxy)perbenzoate (all
15 isomers), 2-ethylhexyl (nonyloxy)perbenzoate (all isomers), Bis (tertbutylmonoperoxy phthaloyl) diperoxy terephthalate, Bis (tertamylmonoperoxy phthaloyl) diperoxy terephthalate diacetyl phthaloyl diperoxide, dibenzoyl phthaloyl diperoxide, bis(4 methylbenzoyl) phthaloyl diperoxide, diacetyl terephthaloyl di peroxide, dibenzoyl terephthaloyl diperoxide and Poly[
dioxycarbonyldioxy(1,1,4,4-tetramethyl-1,4-butanediyl)] peroxide.

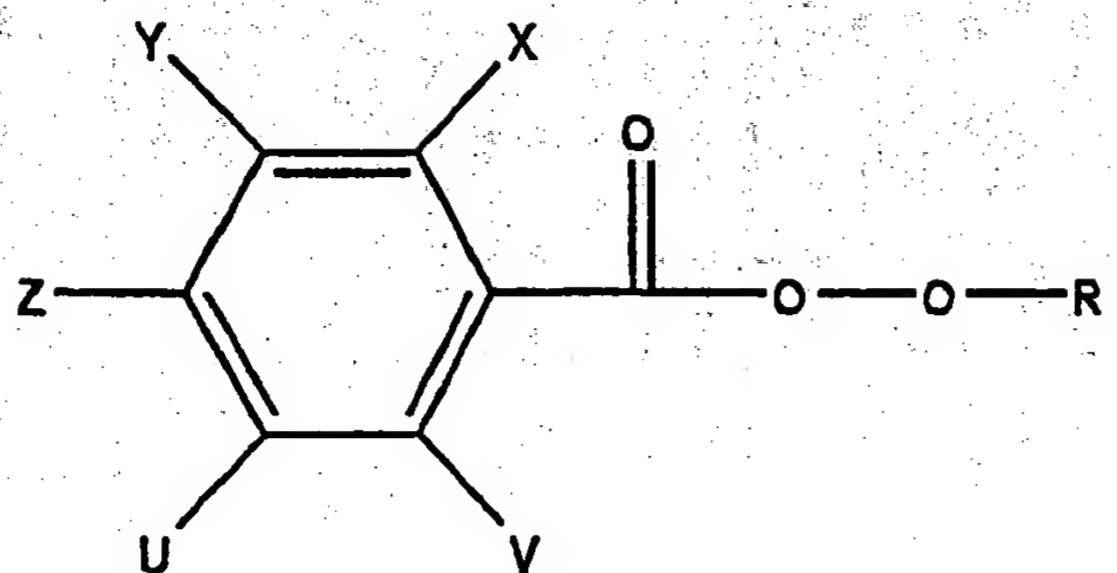
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16. A modified polypropylene produced according to any one of the processes of claims 1, 10 and 12.

17. A process wherein the modified polypropylene of claim 16 is melt mixed with an
25 unmodified polypropylene to produce a modified polypropylene.

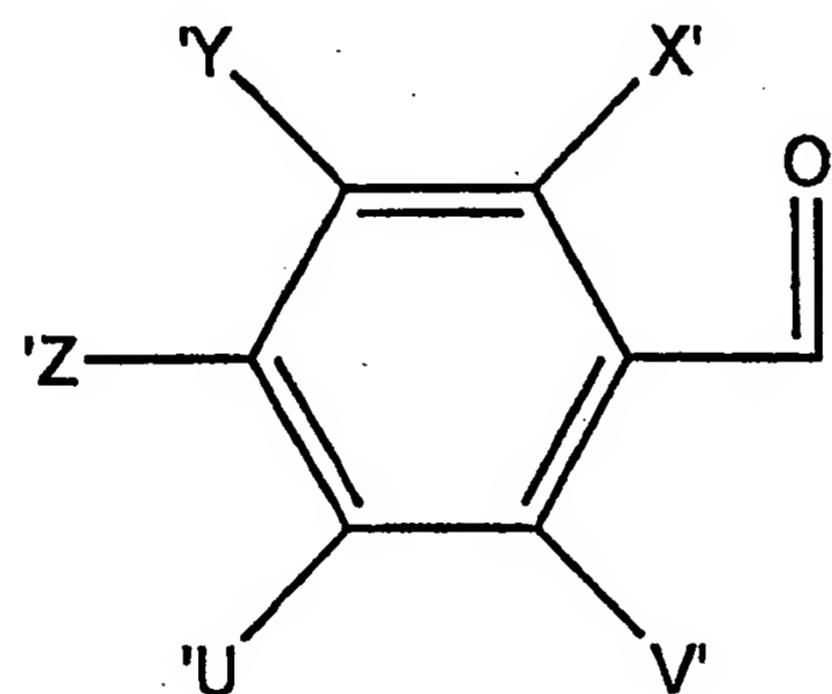
18. A process for modifying a α -olefin polymer wherein said process comprises melt mixing the α -olefin polymer in the presence of an initiator wherein said initiator is selected from the group defined by formula 1.

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Formula 1

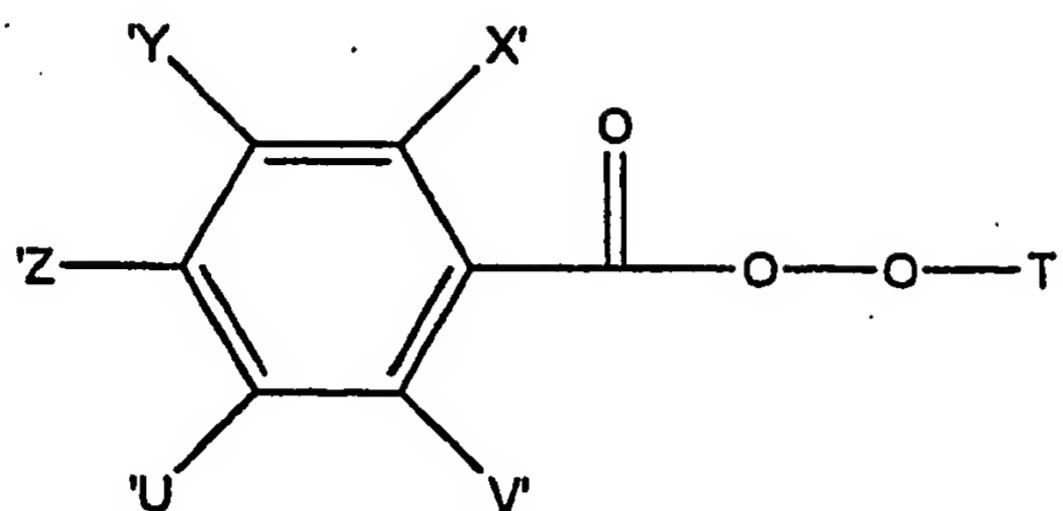
wherein R is selected from the group consisting of optionally substituted C₁ to C₁₈ acyl,
5 optionally substituted C₁ to C₁₈ alkyl, aroyl defined by formula 2,



Formula 2

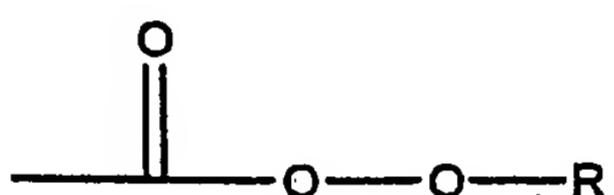
10 and compounds of formula 3,

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Formula 3

wherein U, V, X, Y, Z, U', V', X', Y' and Z' are independently selected from the group
5 consisting hydrogen; halogen; C₁-C₁₈ alkyl; C₁-C₁₈ alkoxy, aryloxy, acyl, acyloxy, aryl,
carboxy, alkoxycarbonyl, aryloxycarbonyl, trialkyl silyl, hydroxy, or a moiety of formula 4,



Formula 4

10 and wherein T is alkylene.

INTERNATIONAL SEARCH REPORT

International application N.
PCT/AU 99/00036

A. CLASSIFICATION OF SUBJECT MATTER

Int Cl⁶: C08K 5/14, C08J 3/24, C08L 23/12, 23/26

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC: as above

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

WPAT & JAPIO: melt() or molten or initiator

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 97/49759 A (AKZO NOVEL N.V.) 31 December 1997 See page 8 lines 15-23, Example 18	1,3,10,16,18
X	EP 0601682 A (ROHM AND HAAS COMPANY) 15 June 1994 See page 6 lines 50-55, Examples 1, 4 and 6	1,4,6,12,15-16,18
X	EP 0528600 A (ROHM AND HAAS CO.) 24 February 1993 See column 4 lines 25-55, Examples 1, 3-5 and claim 9	1,4,6,12, 15-16,18

Further documents are listed in the continuation of Box C

See patent family annex

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier application or patent but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- "&" document member of the same patent family

Date of the actual completion of the international search
8 February 1999

Date of mailing of the international search report
12 FEB 1999

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/AU 99/00036

C (Continuation).

DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	Derwent Abstract accession No. 91-011054/02, Class A17 E17, JP 02283-742 A (TOYADA GOSEI KK) 21 November 1990	1,16,18
X	Derwent Abstract Accession No. 94-313853/39, Class A17, JP 06240096 A (DAI NIPPON INK & CHEM INC) 30 August 1994	1,12,16,18
A	US 5578682 A (WHITE) 26 November 1996 See whole document	1,16,18

INTERNATIONAL SEARCH REPORT
Information on patent family members

International application No.
PCT/AU 99/00036

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report				Patent Family Member			
WO	97/49759	AU	32584/97				
EP	601682	AT	162621	AU	36753/93	BR	9302087
		CA	2093484	DE	69316723	FI	931760
		HU	66372	IL	105359	JP	6207055
		MX	9302288	NO	931357	US	5397842
		ZA	9302747				
EP	528600	AU	20991/92	BR	9203212	CA	2075675
		FI	923712	HU	64091	IL	102757
		JP	5230293	MX	9204774	NO	923174
		US	5397842	ZA	9206053		
US	5578682	CA	2220000	CZ	9703718	EP	828761
		PL	323499	WO	9637521		
END OF ANNEX							

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